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Recent innovations in the technology and applications of low-dimensional CuO nanostructures for sensing, energy and catalysis

Oleg Baranov,^a Kateryna Bazaka,^b Thierry Belmonte,^d Claudia Riccardi,^e H. Eduardo Roman,^e Mandhakini Mohandas,^f Shuyan Xu,^g Uroš Cvelbar^b and Igor Levchenko^h

Low-dimensional copper oxide nanostructures are very promising building blocks for various functional materials targeting high-demanded applications, including energy harvesting and transformation systems, sensing and catalysis. Featuring a very high surface-to-volume ratio and high chemical reactivity, these materials have attracted wide interest from researchers. Currently, extensive research on the fabrication and applications of copper oxide nanostructures ensures the fast progression of this technology. In this article we briefly outline some of the most recent, mostly within the past two years, innovations in well-established fabrication technologies, including oxygen plasma-based methods, self-assembly and electric-field assisted growth, electrospinning and thermal oxidation approaches. Recent progress in several key types of leading-edge applications of CuO nanostructures, mostly for energy, sensing and catalysis, is also reviewed. Besides, we briefly outline and stress novel insights into the effect of various process parameters on the growth of low-dimensional copper oxide nanostructures, such as the heating rate, oxygen flow, and roughness of the substrates. These insights play a key role in establishing links between the structure, properties and performance of the nanomaterials, as well as finding the cost-and-benefit balance for techniques that are capable of fabricating low-dimensional CuO with the desired properties and facilitating their integration into more intricate material architectures and devices without the loss of original properties and function.

1. Introduction

Hierarchical metamaterials (HMMs) endowed with complex architectures and multi-component compositions have drawn much attention in the scientific community.^{1–4} This is because the very existence of designed (intra-material and material–environment) interfaces with specific properties makes HMMs superior to simpler, often single-component, materials. The list of actual applications of HMMs is enormous, with the most relevant ones being biotechnology,^{5,6} sensors^{7–9} and biosensors,^{10,11} energy

harvesting systems,^{12,13} nanomedicine,^{14,15} photocatalysis¹⁶ and chemical catalysis,^{17,18} hydrogen energy,¹⁹ water purification,²⁰ environmental protection²¹ and space exploration.^{22–25}

Metal-oxide nanomaterials

Over the past twenty years, researchers have investigated metal-oxide nanomaterials extensively, showing their adaptivity to different length scales and presenting competitive morphological properties for various applications (Fig. 1a). The benchmark among most of the metal-oxides is represented by copper-oxide, which can form different types of nanostructures, including special topologies such as 1D filament-like structures known as nanowires (NWs). The origin of their unique properties can be found in their extremely high surface-to-volume ratios and the way in which they interact with particular crystal faces. Applications cover a broad range of fields, such as in the sensing of various gases,^{26–28} benzyl alcohol,²⁹ ethanol,³⁰ hydrogen,³¹ glucose,^{32,33} hydrogen sulphide,³⁴ and many other chemicals, as well as in catalysis³⁵ and photocatalysis,^{36–38} batteries,^{39,40} optoelectronics,⁴¹ field emission,⁴² supercapacitor applications,^{43–45}

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broadband visible-light-driven surface-enhanced Raman scattering,^{46,47} and others.^{48–50}

The role of CuO low-dimensional nanomaterials

Similar to many other metal oxide nanostructures, forests of CuO nanowires demonstrate advanced properties that are particularly needed for many applications such as devices for field emission applications,⁵¹ batteries,⁵² supercapacitors,^{53,54} and others. By contrast, many copper oxide-based nanomaterials and nanostructures feature some intrinsic shortcomings, for example, they have relatively low conductivities, low electrochemical stability and high

agglomeration rates. A possible solution for overcoming these shortcomings is to synergistically combine copper oxide-based nanomaterials with other materials such as carbon nanotubes, conducting polymers, and nanomaterials engineered based on graphene hybrids.^{55,56} This can be achieved *via* the utilization of various well-established technologies (see Fig. 1b) including low-temperature chemical synthesis,⁵⁷ high temperature annealing,⁵⁸ the *in situ* modification of various copper materials like copper foam,⁵⁹ anodization in a solution of sodium bicarbonate,⁶⁰ thermal oxidation of copper nanostructures,⁶¹ facile solution-phase treatment techniques,⁶² oxygen oxidation of foils⁶³ and many other techniques.⁶⁴

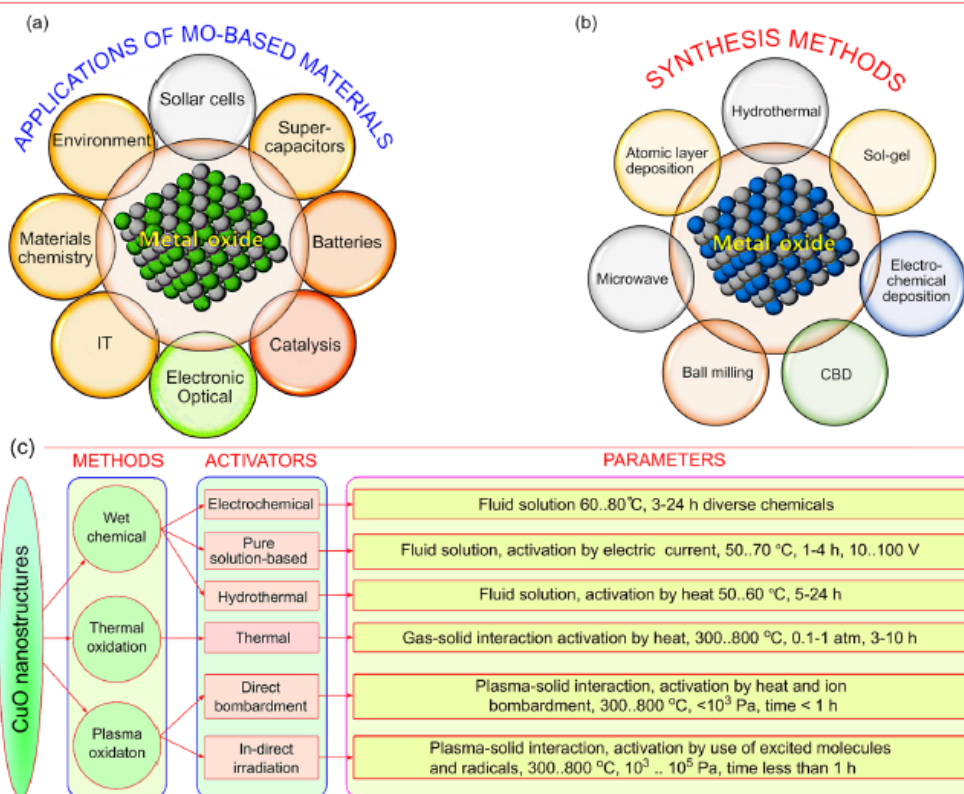


Fig. 1 Spectrum of key technologies and applications of metal oxides, and metal oxide (MO)-based materials. (a) The major key technologies and (b) the most important applications of metal-oxide materials. Virtually all important applications could benefit from the implementation of metal oxide-based materials – from optics and IT to energy harvesting and conservation, catalysis and environmental techniques. Adapted from Ansari *et al.* ref. 65 under conditions of the CC BY license. Copyright 2022. (c) Several key technologies, activators and process parameters specifically for CuO nanomaterials. While the methods can be subdivided into three major groups (wet chemical techniques, thermal and plasma-based methods), the process activators and process parameters demonstrate a flourishing diversity of approaches and conditions, thus offering a great spectrum of possibilities for the future development and sophistication of the MO nanomaterials.

Significance of this review

Featuring very high surface-to-volume ratios and high chemical reactivity, low-dimensional copper oxide nanostructures are very promising building blocks for various functional materials, and thus a review of their properties and potential applications will be of great interest to a wide range of researchers, including, but not limited to the following:

- ✓ the use of low-dimensional copper oxide nanostructures may help overcome the challenges and achieve greater performances for devices used in energy harvesting and transformation, sensing, catalysis and biomedical technologies;

- ✓ copper oxide nanomaterials in general have a very wide spectrum of applications ranging from environmental and biomedical to energy, optics, catalysis and others – moreover, they can be successfully combined with graphene-based materials;

- ✓ finally, in this review article we discuss the very recent progress – mostly within the past two years – in the design of complex, sophisticated low-dimensional copper oxide-based material systems for various applications, aiming to emphasise

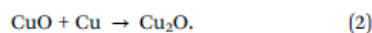
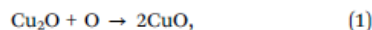
some of the most interesting and important, from the authors' point of view, innovations in the fabrication methods and applications of low-dimensional copper oxide nanostructures.

How are the CuO nanowires formed on the surface?

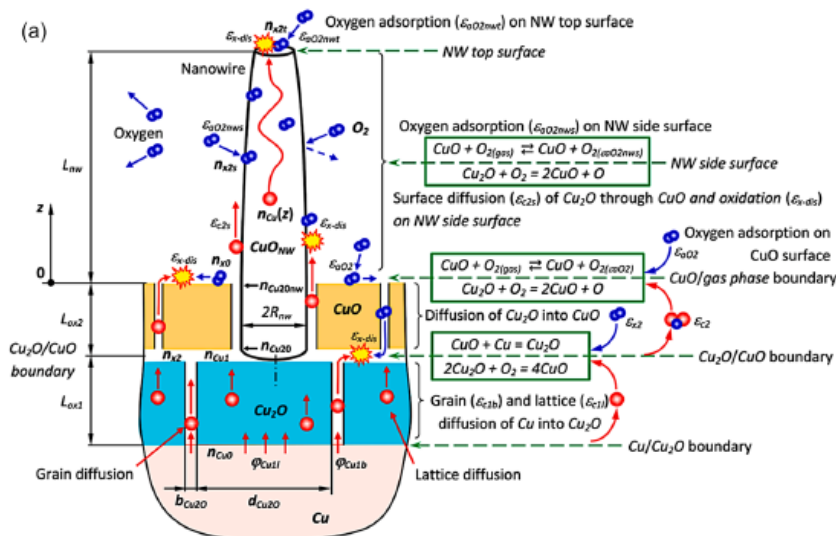
In many fabrication technologies (Fig. 1c), low-dimensional CuO nanostructures are growing directly on substrates. Apart from the convenience of substrate-utilization processes, this approach is suitable for the production of many important

devices, e.g., electrodes for supercapacitors and batteries, catalytic systems, etc., where hierarchical systems consisting of substrates and arrays on nanostructures are required.

Fig. 2a shows a schematic of the thermal growth of nanowires on a substrate as well as the key processes that supply material for the growth. Cu₂O and CuO oxides grow as follows:^{6,6}



SCHEME OF CuO NANOWIRE GROWTH ON SURFACE



DIFFERENT MODES OF COPPER DIFFUSION TOWARD THE NANOWIRE TOP

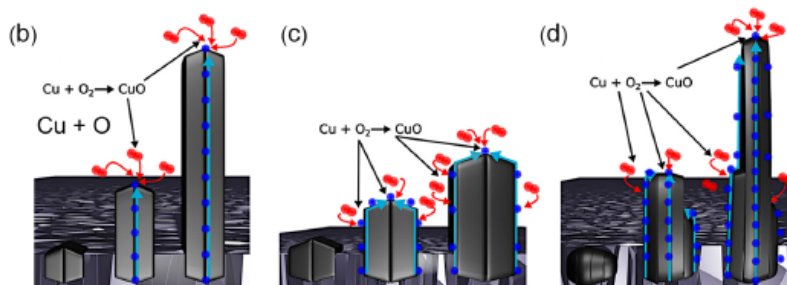


Fig. 2 Schemes and processes of CuO nanowire growth on a surface. (a) Schematic of the thermal growth of nanowires on a substrate and the key processes that supply material for the growth. Reproduced with permission from Baranov et al., ref. 66. Copyright 2021, Elsevier. The gradients of copper and oxygen concentrations on the exposed surfaces of CuO oxide, the boundary between Cu₂O and CuO oxides, and the boundary between Cu and Cu₂O layers are considered as the main driving forces of the process in the formation of CuO nanostructures. Nanowire nuclei are considered as having an anisotropy with respect to the adsorption of oxygen from the gas phase, with the energy of adsorption on the nanowire tips being higher than that on the nanowire side surface; thus, growth on the tips is preferential. Copper atoms are supplied to the surfaces of the nanowires via lattice and grain-boundary diffusion through the Cu₂O layer, grain-boundary diffusion through the CuO layer, and surface or grain-boundary diffusion along the length of the CuO nanowire. The density of surface defects of the side surfaces of the nanowires plays an important role in the formation of long nanowires with a high aspect ratio, since for the perfect surface the oxygen adsorption is low, and copper atoms travel from the nanowire root to the tip without being involved in the reaction of oxidation. As a result, the nanowire does not increase in diameter, and the whole copper flux is delivered to the tip. Opposite to that, the defect-containing side surface promotes oxygen adsorption and increases the rate of oxidation on the side surface of the nanowire; thus, short and bulky nanowires are formed. (b–d) Different modes of copper diffusion toward the nanowire top. Reproduced with permission from Baranov et al., ref. 67. Copyright 2022, ACS.

Fig. 2b–d illustrate the different modes of copper diffusion toward the nanowire top, which can result in quite different morphologies of the resultant nanoarrays of CuO nanowires. SEM images of typical CuO nanowire arrays grown in oxygen at different temperatures, specifically at 600 and 700 °C, demonstrate a significant difference in the nanowire arrays due to the 100 °C difference in temperature, which reveals very complex characteristics of the growth scenario. Specifically, copper is delivered to the growth zones, *viz.*, surface diffusion and lattice diffusion, or both depending on the growth conditions. Copper atoms move *via* jumps, *i.e.*, the atom that is released from the dissociation of the oxide shifts to the nearest node that catches the atom. Detailed models of the growth were recently developed to describe the various morphologies and properties of copper oxide nanowires.⁶⁶

2. Innovations in CuO nanowire fabrication methods

Recently, many advanced techniques for the fabrication of one-dimensional copper oxide nanostructures and arrays have been

developed and tested (Fig. 1). Here we will briefly outline the recent innovations in several of the most important technologies for the fabrication of CuO nanostructures, highlighting the examples that exhibit advanced properties for their applications.

2.1. Recent innovations in plasma-based and electric field-assisted technologies

Among other technologies, plasma-based approaches for growing various nanostructures are very efficient due to the presence of highly energetic ions that efficiently heat and activate the growing nanostructures, and the presence of electric fields that help to form the targeted structures.^{70,71}

Oxygen plasma-based technology. Oxygen plasma for Cu oxidation is efficient since highly energetic ions heat and oxidise the copper atoms. Fig. 3a shows a typical experimental system that can be used for the formation of CuO nanowire arrays on relatively large substrates. This setup is based on a cylindrical (40 mm diameter, 800 mm long) quartz tube where the plasma discharge is sustained. Over the tube, a coil with 9 turns was installed. A radio frequency generator working at 13.56 MHz and 150 W was used to power the coil. The system

RECENT INNOVATIONS IN CuO NANOWIRE FABRICATION METHODS

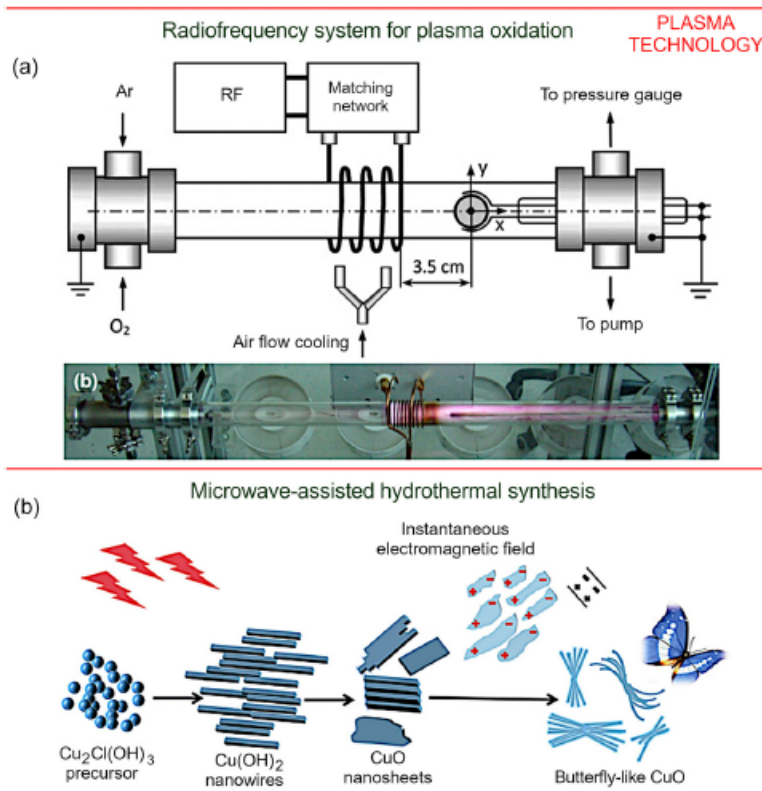


Fig. 3 Plasma-based technology for CuO nanowire fabrication. (a) A typical radio-frequency system for copper oxidation and copper oxide nanowire growth in plasma. The inset show an optical photograph of the setup in operation. Reprinted with permission from Filipic *et al.*, ref. 68. Copyright AIP. (b) Synthesis of butterfly-like copper oxide nanostructures using microwave-assisted hydrothermal technology. Reprinted with permission from Li *et al.*, ref. 69. Copyright Springer Nature, 2020.

also incorporates a vacuum system and pumps, typically a two-stage rotary pump or turbomolecular pumps.

The system used commercial-grade oxygen and argon for the reactive plasma environment. The gas pressure in the tube was monitored using a system of gauges. During the operation, the tube was cooled using a flow of pressurized atmospheric air. The CuO nanowires were formed using copper disc samples, with dimensions of 10 mm diameter and 1 mm thickness, on which the thin, long (4.5 μm) nanowires were successfully fabricated in this system.⁶⁸

Fig. 3b illustrates the synthesis of butterfly-like copper oxidation nanostructures using microwave-assisted hydrothermal technology. The energy carried by the microwaves produces heat which facilitates the reaction, enhancing diffusion of the substances and facilitating their attachment and thereby the growth process. The microwaves directly assisted the intermediate formation of $\text{Cu}(\text{OH})_2$ nanowires, followed by a dehydration process yielding nanosheets of CuO, to produce an associated orthorhombic to monoclinic structural transformation. Due to their polar properties, the CuO sheets were polarized using the microwaves, and arranged finally into a butterfly-like morphology along the external microwave field. It is observed that the size of the CuO butterfly-shaped structures increases with the duration of the applied radiation, suggesting that microwave fields play an essential role in the formation of these special morphologies.⁶⁹

Self-assembly and electric-field assisted growth. Fig. 4a illustrates the self-assembly of CuO nanowires in nanosheets yielding zigzag-shaped twin boundaries. In order to keep the growth of the nanowires, it is necessary to maintain an almost constant oxygen concentration gradient in the cell. Detailed structural analysis suggests that the fast diffusion of ions can help in the self-assembly of single nanowires, yielding the formation of novel twin domain regions characterized by the above-mentioned zigzag-shaped twin boundaries. In addition, such nanosheets displaying high specific surface areas may lead to novel physical properties, e.g., high stiffness, making them suitable as new building blocks for future nanodevices.⁷²

The growth of CuO nanowires, using an electric-field assisted oxidation method, on both sides of a copper foil has recently been reported by Sondors *et al.*⁷³ (Fig. 4b). It was found that increased humidity during the early stages of growth produced higher yields of NWs with diameters smaller than 100 nm, and correspondingly large aspect ratios. Moreover, NWs with diameters smaller than 50 nm possess unusually large Young's moduli in the range of 200–500 GPa. Regarding their electrical properties, the NWs display relatively large electrical resistivities, comparable to those of CuO NWs, and were higher than those obtained when the electric field was applied in wet air. These findings are quite interesting since a combination of low conductivity and high stiffness is excellent for the design of nanoelectromechanical switching devices intended for operation under extreme environmental conditions, such as under high voltages or in harsh environments.

An example of the plasma-based deposition system is shown in the upper panel of Fig. 5.⁷⁴ This system is suitable for the

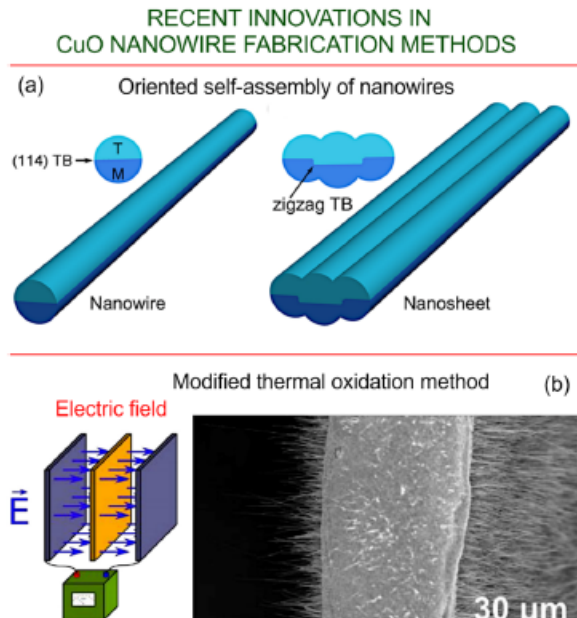


Fig. 4 Self-assembly and electric-field assisted growth. (a) Schematic showing the oriented self-assembly of nanowires, featuring the formation of a zigzag twin boundary (TB). Reprinted with permission from Cao *et al.*, ref. 72. Copyright APS. (b) Growth of CuO nanowires, via a modified thermal oxidation method, on both sides of copper foil using an external electric field. Reprinted from Sondors *et al.* 2020, ref. 73 under terms and conditions of CC BY license.

production of nanostructures with different morphologies (NWs, nanobelts, *etc.*) from various metal oxides including zinc, iron, and copper. Examples of such CuO nanostructures are presented in Fig. 5. Copper oxide NWs can be grown on thin copper foils, yielding structures of different diameters and lengths, depending on the temperature. For example, at 300 $^{\circ}\text{C}$ the mean diameter and length values are about 70 nm and 0.8 μm , respectively, while at 400 $^{\circ}\text{C}$ they increase to 600 nm and 1.4 μm , respectively. The presence of single-crystalline structures associated to form CuO NWs can be revealed using TEM analysis, examples of which are shown in Fig. 5d and e for the case of treatment at 300 $^{\circ}\text{C}$ and of 30 min duration.

Electrospinning. The fabrication of porous SnO_2 -CuO hollow nanofibers using a two-step technology that involves single-needle electrospinning combined with thermal processing to produce nanotubes (NTs) with an augmented specific surface area is illustrated in Fig. 6. The NTs have a wall thickness that is relatively large (15 nm), ensuring a quite homogeneous distribution of copper oxide NPs inside their cage.⁷⁵

The listed examples prove that the synthesis method of the CuO nanostructures greatly affects their performance. Up to now, a number of chemical methods have been developed to synthesize CuO nanostructures. CuO nanowires and nanosheets have been grown directly on copper sheets in alkaline solution *via* single-step voltammetry in the experiments of Singh *et al.*⁷⁶ Kulkarni *et al.*⁷⁷ reported the synthesis of copper oxide nanowires *via* a facile solution-phase method, and the formation of CuO nanowires

RECENT INNOVATIONS IN CuO NANOWIRE FABRICATION METHODS

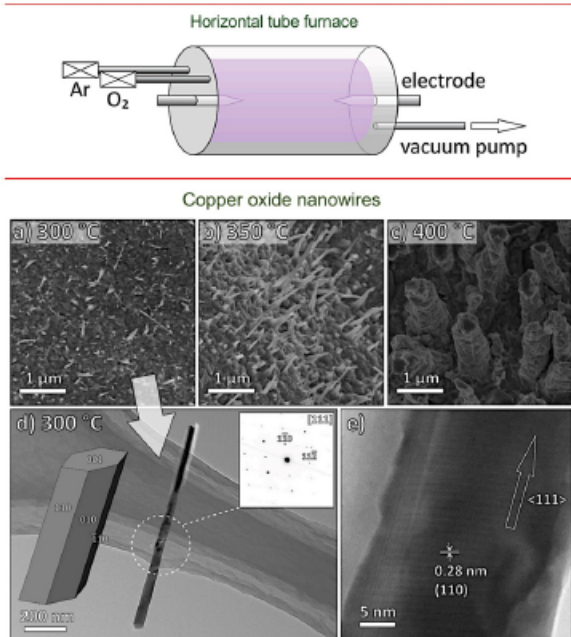


Fig. 5 Plasma-based synthesis of CuO nanowires. Upper panel: schematic of the plasma-assisted furnace deposition chamber contained within a vacuum-sealed glass tube, showing the two pointed-end electrodes that are used to ignite the plasma. Bottom panel: (a–c) SEM images of the copper oxide nanostructures synthesized at different growth temperatures for growth time of 30 min; (d) TEM image of a single copper oxide nanostructure grown at 300 °C viewed along the [111] zone axis and a nanowire model in the same orientation (inset shows the selected-area electron diffraction pattern); and (e) high-resolution TEM image of the copper oxide nanostructure with the marked interplanar spacing. Reprinted from Guo *et al.*, ref. 74 under terms and conditions of CC BY license.

obtained *via* anodization in sodium bicarbonate solution was studied by Gizinski *et al.*⁷⁸ Other chemical^{79,80} or plasma-enhanced⁸¹

methods have also been developed and have the advantages of being controlled easily through involving the presence of reagents, or electric and magnetic fields.^{82,83}

2.2. Recent innovations in thermal and wet-chemical technologies

Thermal oxidation. Thermal oxidation is one of the most widely used technologies for the fabrication of CuO nanowires. Not surprisingly, it is currently under intense investigation. CuO nanowires have been synthesized successfully *via* thermal oxidation in air by Deng *et al.*⁸⁴ (Fig. 7). They found that increasing the annealing temperature and treatment time resulted in an increased density of NWs. Specifically, by increasing the temperature in the range from 400 to 600 °C it was found that the NW diameter varied from 140–160 nm to 330–380 nm, respectively. However, an increased NW length was favored when both the annealing temperature and time were increased. No NWs formed below the 300 °C annealing temperature, but above 400 °C a few short NWs start to be seen. At higher temperatures, say between 500 °C and 600 °C, a large amount of densely packed, long NWs emerge, where a steady diameter increase is seen with the annealing temperature.

The thermal oxidation results discussed so far are indissolubly linked to a diffusion process of CuO along grain boundaries, in which a Cu₂O layer is formed that acts as the precursor for the incipient NWs. Generally, the length of the nanowires increases with the annealing time. For example, the average length of 5.1 μm was obtained in the case of 2 h of treatment, which increased to 10.2 μm after 8 h of treatment, confirming the effect of the annealing time on the length. By contrast, the NW diameter turns out to be almost independent of the treatment time. This is attributed to the fact that the NW diameter is highly dependent on the grain size, which determines the cross-section geometry of the wires, but which is not sensitive to thermal annealing.

RECENT INNOVATIONS IN CuO NANOSTRUCTURE FABRICATION METHODS

Fabrication of porous SnO₂-CuO hollow nanofibers

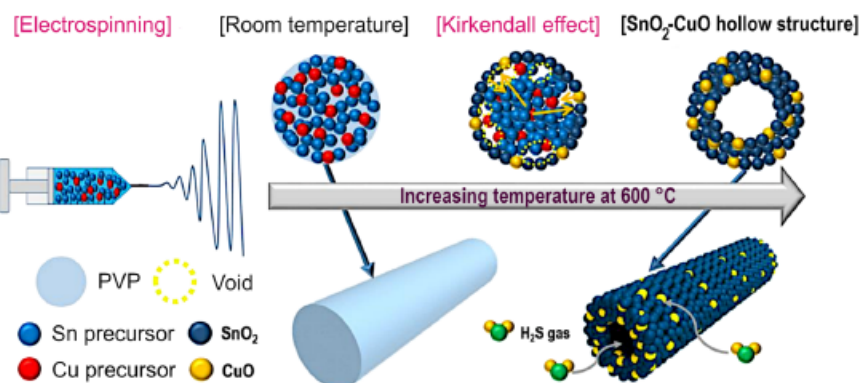


Fig. 6 Electrospinning combined with thermal processing. The fabrication of porous SnO₂-CuO hollow nanofibers for gas sensors *via* electrospinning combined with thermal processing. Reprinted from Park *et al.* 2020, ref. 75 under terms and conditions of CC BY license.

Nanowires grown directly on V-shaped microgrooves

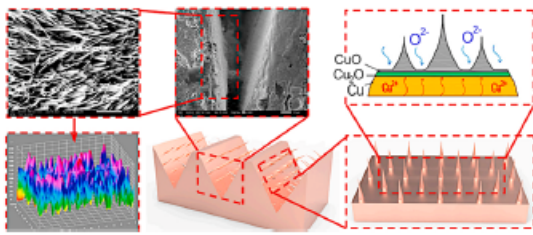


Fig. 7 Nanowires grown directly on V-shaped microgrooves. After thermal oxidation, the nanowires were grown directly on V-shaped microgrooves. Importantly, it was found that both microvoids and microcavities appear to play an essential role in the process. Reprinted with permission from Deng *et al.*, ref. 84. Copyright 2021, Elsevier.

The results obtained by Jafari *et al.*⁸⁶ support the above-mentioned conclusions, although at higher temperatures, in the range of 500–700 °C. At 700 °C, nanowires form denser arrays reaching 1–3 μm in length, thus suggesting that compressive stress is important for determining both the nanowire diameter and density. They show remarkable photocatalytic properties and electrical conductivity, which could be associated with different electron scattering mechanisms. This general picture was also confirmed by Mahmoodi *et al.*,⁸⁷ who performed experiments at the lower temperature of 400 °C, in which the NW length increased from about 1 μm after 5 h to 2–3 μm after 5–7 h, and to 2.5–3.5 μm after 7–9 h of annealing time. As expected, the NW diameter was insensitive to the treatment time, falling within the range of 50–70 nm.

Experiments reported by Feng *et al.*⁸⁸ were performed at three temperatures (400, 450, and 500 °C) and at treatment durations ranging from 4 to 12 hours. They confirmed that both the diameter and the length of the nanowires increased with temperature and the growth time. Differences in aspect ratio trends were observed when varying either the temperature at a fixed time of growth or the growth time at a fixed temperature. Interestingly, a maximum diameter was found at around 8 h for each temperature considered.

Interesting technology has been recently reported by Sun *et al.*⁸⁵ A Cu wafer was used as the source of copper for the initial synthesis of Cu₂S nanoparticles. During the oxygen evolution reaction, sub-micrometer Cu₂S particles transformed into nanowire-like CuO nanostructures (Fig. 8).

Thermal oxidation was used recently to fabricate detectors with a high level of sensitivity, as well as a fast and stable photoresponse to visible light, using well aligned copper oxide nanowires.⁹⁶ The absorption of visible light was increased substantially by exploiting the interference effects that occur in nanowire arrays of certain geometry. The desired combination of geometry and alignment was realized through the use of an SiO₂ template that was fabricated using repeated cycles of wet-chemical oxidation and etching, as they enable the generation of 3D arrays of regularly spaced, well-aligned nanoscale structures on larger surfaces with high (1 nm) resolution (Fig. 9b). The glancing angle deposition (GLAD) technique was then used to

Cuprous sulfide derived CuO nanowires

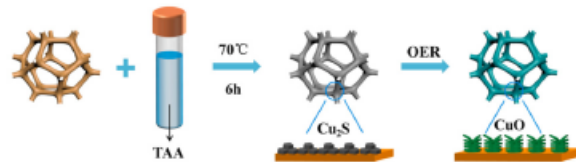


Fig. 8 Cuprous sulfide derived CuO nanowires. Synthesis of Cu₂S/CF and the related phase change during operation of the oxygen evolution reaction (OER). Reprinted with permission from Sun *et al.*, ref. 85. Copyright 2021, Elsevier.

grow copper nanowires on the surface of the thus-fabricated SiO₂ template, with high-temperature oxidation at 700 °C for 3 h in air then used to convert Cu into CuO. Next, gold electrodes were deposited by first coating the surface of the CuO nanowires with a photoresist *via* spin coating, growing a layer of Au using PVD, and then removing the photoresist. SEM images of the obtained structure and schematics of the device are shown in Fig. 9.

Wet-chemical and hydrothermal technologies. Wet-chemical and hydrothermal routes for CuO nanomaterial fabrication are among the most important techniques that are currently under development for critical applications such as in lithium-ion batteries, where wet-chemical methods ensure the fabrication of, for example, electrode materials,^{89–91} optical materials,⁹² photocatalysts,⁹³ bioactive materials⁹⁴ and other functional nanomaterials.⁹⁵

Fig. 10a illustrates an interesting example of the application of the wet-chemical process to fabricate ultrathin CuO nanowires using self-assembled Cu(OH)₂ as the starting material.⁹⁷ Cu(OH)₂ NW structures are fabricated from an inorganic colloidal suspension of Cu(OH)₂, where the selective adhesion of organic solvent molecules changes the surface energy of the inorganic crystal, which in turn drives the aggregation of crystallites and, from a practical standpoint, enables the fabrication of aggregates with the desired size and shape. In this particular study, such directed crystallite growth was used to produce Cu(OH)₂ NWs with an average aspect ratio of > 112. Subsequent annealing of the thus-produced Cu(OH)₂ NWs resulted in the formation of CuO nanowires, where the length of the nanowire remained similar (~7 μm), and the average diameter of the CuO nanowires increased six-fold, to ~27 nm. This simple approach provides a potentially useful avenue for the fabrication of electronically-active nanostructures with the desired size and shape configurations.

Another interesting material that is currently being investigated is the so-called ‘hybrid’ CuO/Cu₂O nanowires, for which the mechanism of formation is illustrated in Fig. 10b.⁹⁸ The CuO/Cu₂O nanowires were synthesized directly from copper acetate, which is both abundant and affordable, using wet chemistry. This very simple process consisted of mixing the copper acetate solution in double-distilled water with ammonia solution, followed by thorough stirring and the addition of sodium hydroxide. The synthesis was conducted at 90° for two hours, with stirring. After keeping the solution overnight at room temperature, precipitates that had formed were centrifuged,

RECENT INNOVATIONS IN CuO NANOWIRE FABRICATION METHODS

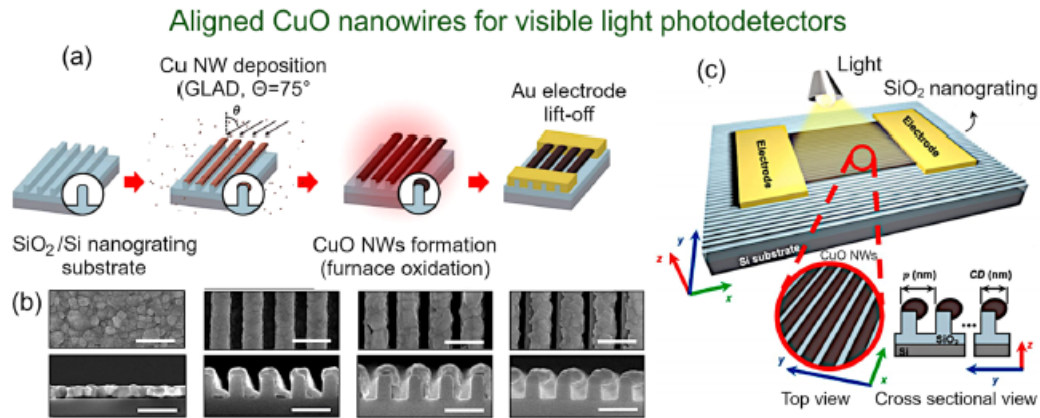


Fig. 9 Aligned CuO nanowires for visible-light photodetectors via thermal oxidation. (a) Method for the growth of well-aligned CuO nanowires using a combination of substrate nanograting, GLAD and furnace oxidation, with the subsequent deposition of Au electrodes using a combination of photoresist lithography and physical vapor deposition. (b) SEM images of the copper oxide film and nanowires with critical dimensions. Top views are shown in the upper panels, and cross-section views are shown in the bottom panels. The length scale is 500 nm. (c) Structure of the photodetector illustrating the placement of copper oxide nanowires on the surface of the patterned SiO₂. Reprinted under terms and conditions of CC BY license from Jo *et al.*, ref. 96. Copyright 2022, the Authors.

washed with distilled water and ethanol, and dried. Despite the remarkable simplicity and low price of this technology, the resultant material has demonstrated an extremely high efficiency when used as a catalytic platform for the effective conversion of 4-nitrophenol to 4-aminophenol, the former being a phenolic

compound that is used widely in the fabrication of pesticides, drugs, and synthetic dyes, and which is known to be harmful. The conversion of 4-nitrophenol to 4-aminophenol is a reaction that is also commonly used in the synthesis of analgesics and antipyretic pharmaceuticals, including paracetamol.

RECENT INNOVATIONS IN CuO NANOSTRUCTURE FABRICATION METHODS

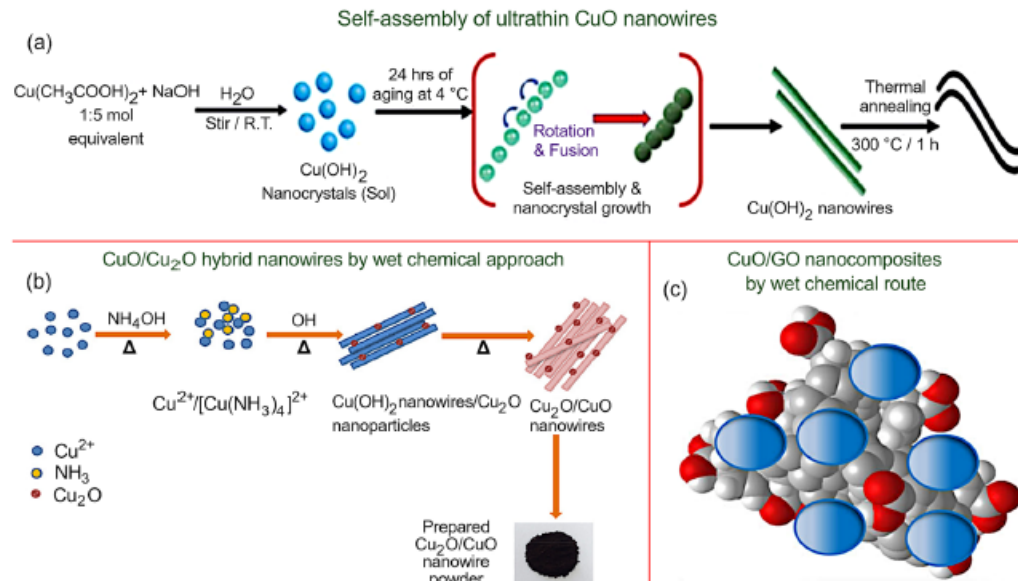


Fig. 10 Wet-chemical and hydrothermal technologies. (a) Illustration of the chemical reaction pathway and the subsequent self-directed growth of Cu(OH)₂ colloidal particles, followed by their thermal conversion to copper oxide nanowires. Reprinted under terms and conditions of CC BY license from Pathiraja *et al.*, ref. 97. Copyright 2020, the Authors. (b) Likely pathway for the formation of hybrid CuO/Cu₂O nanowires. Reprinted with permission from Sahu *et al.*, ref. 98. Copyright 2020, Elsevier. (c) CuO/GO nanocomposites produced via wet-chemical synthesis. Reprinted under terms and conditions of CC BY license from Biswas *et al.*, ref. 99. Copyright 2021, the Authors.

The efficiency with which the CuO/Cu₂O nanowires catalyzed the conversion reaction emphasizes the potential of wet-chemical processes for the nano-synthesis of advanced nanomaterials.

Wet chemical routes also enable the fabrication of very important and promising copper oxide-graphene oxide (CuO/GO) nanocomposite materials. In a recently reported study, a bottom-up approach was used for the synthesis of copper oxide nanoparticles directly on the surface of graphene oxide flakes, the latter being a hydrophilic material. The resulting composites of copper oxide and graphene oxide were obtained by combining two specific precursors, copper nitrate and citric acid, intermittently mixed with graphene oxide solutions.⁹⁹ Importantly, the formation of the C=C, C-O, and Cu-C functional groups was confirmed using FTIR analysis. Morphologically, the composite material consists of CuO nanoparticles deposited on planar hydrophilic graphene oxide sheets (Fig. 10c). These results suggest the possibility of synthesizing nanomaterial-based devices with a high degree of sensitivity that can be used simultaneously for diagnostics and therapy, leading to the development of new solutions in healthcare.

The top panel of Fig. 11 illustrates one of the ways by which petal-shaped nanoparticles of copper oxide can be grown using hydrothermal technology. The thus-obtained petal-shaped monoclinic nanocrystals display a high size uniformity, a considerable surface area and desirable morphological characteristics. Importantly, hydrothermal technology appears to be well suited for the synthesis of petal-like copper oxide nanostructures due to its relative simplicity and affordability, the purity of the resulting products, the high reproducibility of the results, the lower defect levels in comparison with other methods that require high temperatures, and, finally, its remarkable ability to be scaled up for the fabrication of monodisperse copper oxide nanostructures in larger quantities.¹⁰⁰ To fabricate the CuO nanopetals, powdered copper nitrate is first combined with water to produce a uniform dispersion, which is then introduced into an aqueous solution of hexamethylenetetramine at ambient temperature with thorough stirring. Next, the thus-obtained solution is heated to 110 °C for about 3 h under elevated pressure. Finally, it is cooled to ambient temperature, followed by centrifugation and washing. The black solid material produced as a result of this process is then subjected to drying under a nitrogen flow in a vacuum oven. Thus, this process is also very simple, cheap, and does not demand the use of complex and/or expensive equipment. Nevertheless, the produced CuO nano-petals demonstrate an advanced (above 99% in 90 min) catalytic activity for the degradation of methylene blue in the presence of hydrogen peroxide.

Zinc oxide-doped copper oxide (ZnO-CuO) nanoleaves for the non-enzymatic detection of acetylcholine and ascorbic acid were synthesized using a facile wet-chemical technology (Fig. 11, lower panel). Cheap reagents (*i.e.*, zinc chloride, copper sulfate and NaOH) were used. Zinc chloride and copper sulfate were dissolved in water, stirred, and the pH of the solutions was adjusted using NaOH. Next, the solutions were stirred for 6 hours at 90 °C. After evaporation and drying, the material

was ground to a powder.¹⁰¹ The prototype sensor fabricated using this material is capable of detecting acetylcholine and ascorbic acid in samples of human, mouse, and rabbit serum, as well as in orange juice and urine.

Wet-chemical technology was also used to produce urchin-like cobalt-doped copper oxide nanostructures for antimicrobial applications and the removal of organic pollutants from water.¹⁰² The Co-doped CuO sample was prepared using relatively cheap reagents such as cobalt nitrate hexahydrate, liquid ammonia and ethanol *via* wet co-precipitation technology. Cobalt and copper salt solutions were mixed and then agitated for 20 min at 50 °C, where stirring if the mixture was continued until precipitation of the residue was achieved, before drying (Fig. 12). Finally, the produced Co_xCu_{1-x}O was chemically activated using KOH. Despite its simplicity and affordability, this approach of combined Co doping and activation has the potential to bring about a notable increase in the performance of copper oxide as a catalytic platform in under visible light.

The preparation of binary metal doped CuO nanocatalysts using liquid reagents is one more example of the wet chemistry application to produce advanced nanomaterials. Doping CuO nanoparticles with cerium and zinc enabled the fabrication of the CeZn-CuO photocatalyst with superior activity.¹⁰⁵ In this process, liquid ammonia was used, simultaneously, as the precipitating agent and pH regulator. This simple co-precipitation technology involving the mixing and stirring of respective hydrated nitrate solutions of cerium, zinc and copper, followed by the addition of ammonia solution as the precipitation agent, resulted in the formation of a nanomaterial, which was then washed and dried, before being ground and annealed at 750 °C (Fig. 13). CeZn-CuO was able to degrade 81.64% MB dye in 100 min.

2.3. Effect of process parameters: recent insights

Effect of the heating rate. Nkhaili *et al.*¹⁰³ performed experiments to elucidate the effects of varying the heating rate on the shape of the NWs grown. CuO NWs were grown *via* thermal oxidation at 550 °C in air and for 4 h, using heating rates of 1, 2, 5, and 10 °C min⁻¹. The resulting NWs had a similar diameter of about 200 nm when grown on a copper foil of 0.3 mm in thickness. They found that the NW density decreases with the heating rate. Tran *et al.*¹⁰⁴ reported measurements at 400 °C and 600 °C. For the former, the NW length was 3–5 μm, and width was about 30–60 nm. By varying the temperature in the above range, the average NW diameter increased from 30 nm to 200 nm, and the NWs presented a cone-like structure with a smaller diameter towards the tip. In concomitance with the diameter variation, they observed an increment in the grain size of the CuO layer. Based on XRD analysis, the authors explained the dependence of the NW size and density on the grain size, concluding that two mechanisms are responsible for the observed density of the NWs, that is, an efficient diffusion of Cu ions along the enhanced grain boundaries in the oxide layer, which was further increased as the temperature was increased. They also found that larger NWs and lower densities are favoured at high temperatures.

RECENT INNOVATIONS IN CuO NANOSTRUCTURE FABRICATION METHODS

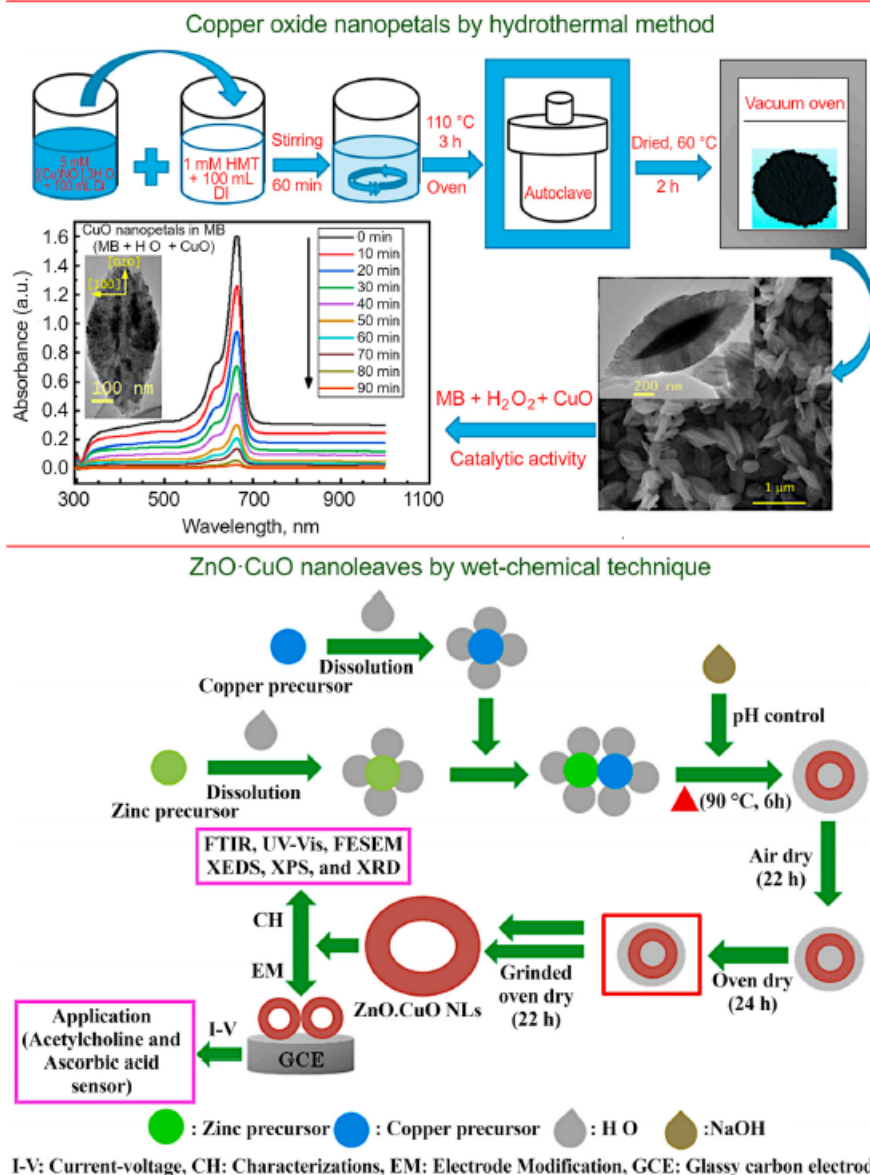


Fig. 11 Wet-chemical and hydrothermal technologies. (Top panel) Petal-shaped CuO nanostructures grown using hydrothermal technology, and their application in catalysis (MB denotes methylene blue). Reprinted under terms and conditions of CC BY license from Khan *et al.*, ref. 100. Copyright 2020, the Authors. (Bottom panel) Synthesis of sensors for the detection of acetylcholine and ascorbic acid using zinc oxide/copper oxide leaf-shaped nanostructures fabricated using wet chemistry. Reprinted with permission from Hussain *et al.*, ref. 101. Copyright 2020, Elsevier.

Zúñiga *et al.* synthesized NWs of 0.3 mm in diameter at 400 °C and 500 °C in air for about 2 h.¹⁰⁶ At the two temperatures, both the diameter and length of the NWs were found to increase with time, and the NWs were found to be composed of bi-crystals. Mohamed *et al.* investigated NW formation from Cu₂O/CuO composites *via* the thermal oxidation of 0.51 mm thick copper foil at 550 °C, 600 °C, and 650 °C, employing as the input an Ar (100 sccm) and O₂ (50 sccm) gas mixture for about 4 h.¹⁰⁷

The NW diameter and length results for the three temperatures considered (550, 600 and 650 °C) were as follows: 95–250 nm, 9 μm; 105–295 nm, 12 μm; and 100–295 nm, 30 μm, respectively. It was found that at 650 °C, some of the NWs break down when their length exceeds a certain value. This was thought to be due to a mismatch between the Cu₂O and Cu lattices, producing a larger amount of stress, indicating that the release of such a significant stress is the actual driving force for the NW growth. A model was proposed by Zhu *et al.*¹⁰⁸ to

RECENT INNOVATIONS IN CuO NANOSTRUCTURE FABRICATION METHODS

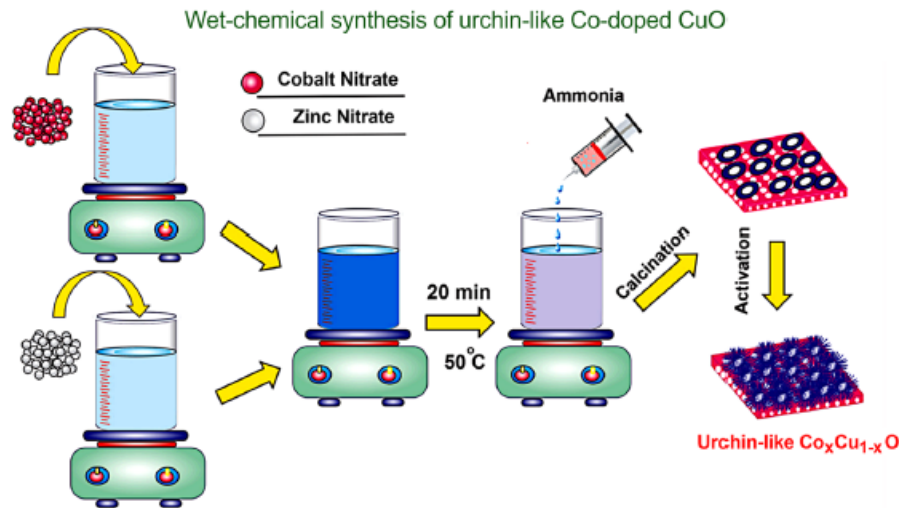


Fig. 12 Wet-chemical and hydrothermal technologies. Fabrication of $\text{Co}_x\text{-Cu}_{1-x}\text{O}$ nanostructures with an urchin-like morphology. The Co-doped CuO nanostructures were prepared using cheap reagents such as cobalt nitrate hexahydrate, liquid ammonia and ethanol via wet co-precipitation technology. Reprinted with permission from Alburai et al., ref. 102. Copyright 2022, Elsevier.

explain how NW bi-crystals can be formed, leaving open the question about mono-crystalline NWs, as mentioned by Zhang *et al.*¹⁰⁹ Besides, two main mechanisms for copper diffusion to the NW top have drawn most attention in the literature:¹¹⁰ one is the diffusion *via* a twin boundary, and the second is just the

surface diffusion of Cu ions on the NW surface itself. Moreover, important results were recently reported by Hu *et al.* who grew CuO nanoneedle arrays on top of commercial ceramic tubes using magnetron sputtering,^{111,112} followed by both wet chemical etching and annealing.¹¹³ Fig. 14 (upper panel) illustrates the

RECENT INNOVATIONS IN CuO NANOSTRUCTURE FABRICATION METHODS

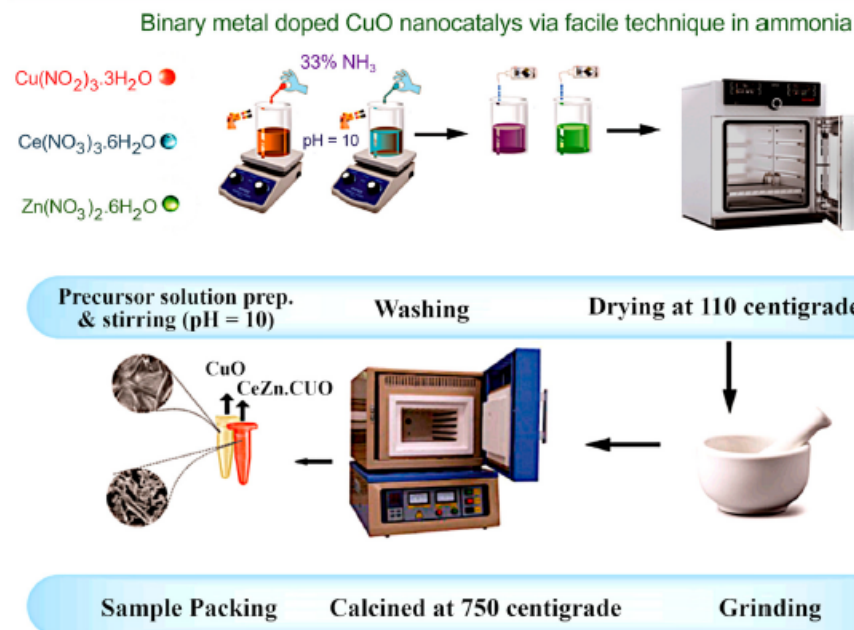


Fig. 13 Wet-chemical and hydrothermal technologies. Synthesis and assembly of CuO nanocatalysts doped with a binary metal. Doping CuO nanoparticles with cerium and zinc enabled the fabrication of CeZn-CuO photocatalyst with superior activity. Liquid ammonia was used as a precipitating agent and pH regulator, simultaneously. Reprinted with permission from Rehman *et al.*, ref. 105. Copyright 2021, Elsevier.

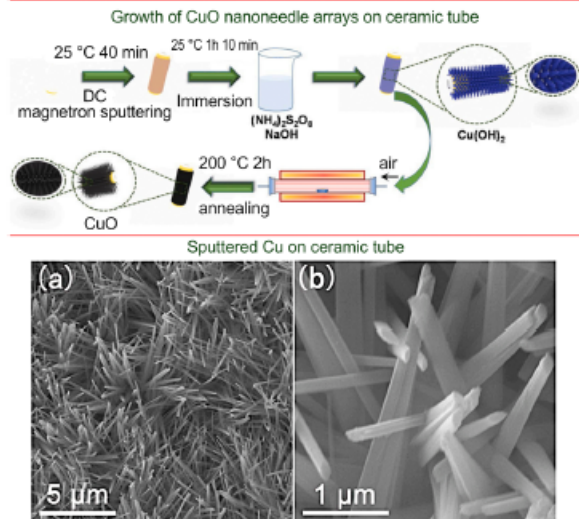


Fig. 14 (Upper panel) Schematic of CuO nanoneedle arrays grown on a commercial ceramic tube. (Lower panel) SEM images at low (a) and high magnification (b) of sputtered Cu on the ceramic tube after wet etching. The nanoarrays are able to keep their morphology after a calcination process at 200 °C. Reprinted with permission from Hu *et al.*, ref. 113. Copyright 2021, Elsevier.

scheme of the process, and Fig. 14 (lower panel) shows the SEM images of the obtained nanostructures. This technology opens the way to the fabrication of cheap tube-like gas sensors.

Effect of oxygen flow. The oxygen flow is one of the important parameters that controls the growth of CuO nanostructures. Not surprisingly, many studies have been devoted to the effect of oxygen flow on the CuO parameters. As shown by Hansen *et al.*,¹¹⁴ the diameter and length of the NWs increase with the oxygen flow and concentration. With heating at 500 °C and oxidation for about 150 min, they considered flow values of 0.5, 0.75, 1.0, and 5.0 mL min⁻¹. From measurements at 5 mL min⁻¹, where the O₂/Ar ratio in the used mixture was varied in the range of 10 to 100%, they concluded that the O₂ concentration is the parameter that actually controls the process. Kumar *et al.*¹¹⁵ also report studies on the effects of oxygen flow, suggesting the existence of a possible optimal flow rate to oxidise the copper surface. Oxidation measurements were carried out at 500 °C, and the flow rates were chosen to be 35, 150, and 300 mL min⁻¹, yielding average NW lengths in the ranges of 1–3 μm, 2–5 μm and <2 μm, respectively. At an even higher flow of 500 mL min⁻¹, no NWs were found. A possible explanation for this behaviour is the absence of seeds for the NWs as a result of an alteration of the stress conditions in the oxide layer. The possible existence of an optimum oxygen flow for NW growth has also been reported recently.⁷² Another interesting effect to be taken into account is that of the humidity, as studied by Xu *et al.*,¹¹⁶ in which they considered wet air and wet oxygen. They concluded that the nanowire morphology is determined to a large extent by the partial pressure of oxygen, while water vapor has opposite effects in air with respect to the oxygen atmosphere. In wet air, the NWs

are denser and more uniform, yielding diameters of about 80 nm and lengths of 3 μm, while in a wet O₂ atmosphere, they have lower densities. The conclusion is that water vapour can affect both the nucleation and the growth rate of the nanowires. Indeed, Sondors *et al.* reported that the NW growth can be affected by the humidity and by the application of an external field.⁷³

Initial roughness of the copper substrate. The effect of the substrate thickness on the growth of NWs has been studied at 400 °C,¹¹⁷ and it was found that no NWs grew on 500 nm thick films, although they appear at the 1 μm level. Rough copper substrates yield smaller grains within the oxide layers, thus favouring grain boundary diffusion,^{118,119} and at the same time leading to higher NW densities. Copper is delivered *via* the Cu₂O layer to the Cu₂O/CuO oxide boundary, thus both lattice¹²⁰ and grain boundary¹²¹ diffusion mechanisms can be present. The former dominates in the range of 350–500 °C whereas the latter dominates at higher temperatures of 800–900 °C,¹²² and both mechanisms will be operating in the model when the NW growth temperature lies between the mentioned intervals. Tang *et al.*¹²³ have shown that impurities and/or cracks can also influence the diffusion process, the phenomenology of which was explained by invoking the Kirkendall effect.¹²⁴ Diffusion results from stress gradients along its path, associated with the different molar volumes belonging to both oxides and the metal phase.¹²⁵ Mema *et al.* succeeded in controlling the mechanical stress generated by the deformation of copper foil, resulting in enhanced NW growth.¹²⁶ Apparently, the macroscopic stresses so generated appear to be of a much lower magnitude than those mentioned above. Finally, Park *et al.* suggested an explanation based on a two-step process yielding the formation of so-called hillocks of copper in order to relieve the stress.¹²⁷ This leads to the formation of a Cu₂O phase through oxidation of the hillocks.

Hybrid systems based on Cu_xO nanowires. Hybrid systems possess great potential for various applications but pose a hard problem regarding their synthesis. Le *et al.* succeeded in the production of a hybrid system based on gold nanoparticles and a Cu_xO nanowire–3D copper foam.¹²⁸ First, the 3D copper foam (3DF) was cut into slices, cleaned, washed and then immersed, at room temperature, into a mixture of 2.5 M NaOH and 0.13 M (NH₄)₂S₂O₈, with gentle stirring for about 15 min. Then, Cu(OH)₂ nanoneedle arrays were grown on the 3DF, and as a result the sample was washed with water and dried. Next, the Cu(OH)₂ NWs–3DF were reduced to Cu NWs–3DF at 200 °C under a mixed flow of Ar and H₂ gas for 80 min. After the reaction had finished, the system underwent rapid cooling to room temperature under the action of an Ar flow (100 sccm); it was then removed and followed by rapid immersion in a 1 mM HAuCl₄ solution for 1 min. This allowed the electroless deposition of the sample, as shown in Fig. 15a. Thus, the resulting Au–Cu@Cu_xO NWs–3DF was carefully rinsed, followed by drying under a flow of N₂ gas. Fig. 15b shows images of the synthesized Au–Cu@Cu_xO NWs–3D foam hybrid structure.

2.4. Recent insights into the driving forces for one-dimensional growth

The question of the driving force for the 1D formation of long nanostructures is still under debate.¹²⁹ Among the possible NW

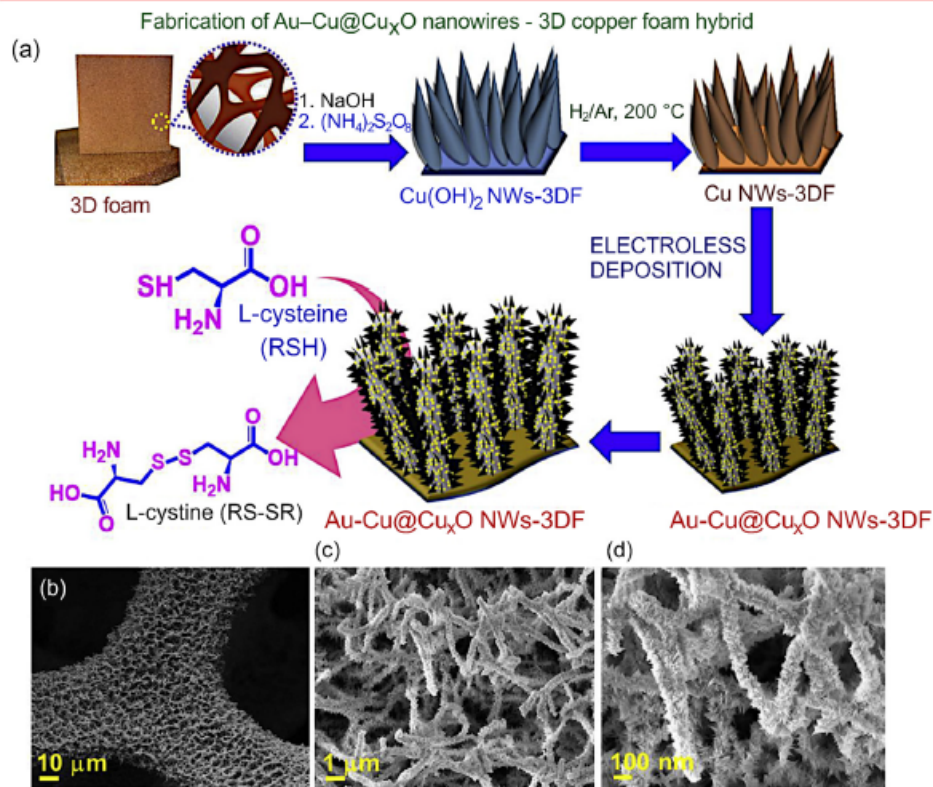


Fig. 15 Hybrid system based on gold nanoparticles and a Cu_xO nanowire-3D copper foam. (a) Schematic of the fabrication of the Au-Cu@Cu_xO nanowire-3D copper foam hybrid. (b-d) SEM images of the synthesized Au-Cu@Cu_xO nanowire-3D foam hybrid structure. Reprinted with permission from Le *et al.*, ref. 128. Copyright 2019, Elsevier.

growth mechanisms, vapour-solid (VS) and vapour-liquid-solid (VLS) approaches are not appropriate for temperatures in the range of 300–800 °C, which are well below the melting point of Cu or its oxide compounds. Recent experiments have suggested that CuO NWs can grow on their tops,^{130,131} suggesting that Cu should be supplied to the tops of nanowires, thus enabling their reaction with oxygen. As we already mentioned above, the possibilities of this can occur *via* Cu diffusion either along the grain boundaries or on the NW surface.¹³¹ However, both explanations have some drawbacks. For instance, in the case of surface diffusion, one would expect thicker NWs at the bottom, while most of the available results point to a uniform NW width distribution. If the diffusion along the grain boundary is assumed, one cannot explain Cu diffusion for cases in which grain boundaries are not present, which have been indeed observed in experiments. Interesting results were obtained by Chen *et al.* with respect to the effect of the annealing time: a large number of long thin nanowires were grown at 500 °C and 8 hours of exposure, and only CuO crystallites were developed at the same temperature but for 24 hours of annealing.¹³² The results are a possible manifestation of the fact that the reaction between oxygen and copper takes place at the side of the NW surface or on its defects,

without the need to invoke any of the above-mentioned mechanisms or even in synergy with them.

Diffusion and synergy as important driving forces for the growth. By measuring the diameter and length of nanowires as a function of time, Chen *et al.* demonstrated that the diffusion of copper along the sides of growing nanowire surfaces led to the formation of oxides along their diffusion paths.¹³¹ Indeed, TEM analysis has shown that above a certain oxygen pressure, twin boundaries act as nucleation sites of newly favoured phases.¹³³ This corresponds with several observations of bi-crystalline nanowire structures displaying twin boundaries in the middle of the surface.^{134,135} Interestingly, Shi *et al.*¹²⁴ have suggested that synergetic effects associated with the thermal growth of CuO NWs are able to explain many of the observed single/bi-crystals and cone-shaped NW structures. Furthermore, Tu *et al.*¹³⁶ obtained single-crystal NWs at 400 °C, while at higher temperatures of 500–600 °C bi-crystal NWs were found.

Conductive films to intensify the growth. A study of annealing and the activation layer for the growth of CuO NWs has been discussed by Chawla *et al.*,¹³⁷ where copper films could be synthesized on silicon-based buffers, using sputtering techniques. The obtained samples underwent a careful annealing process by varying the temperature in the range of 300–700 °C,

for a sufficiently long time between 2 and 8 hours. This procedure enabled the authors to gain insight into the growth mechanism involved in the formation of the NWs. In addition, it was found that the use of a thin gold film on the top of the structure enhanced both the density and the aspect ratio of the NWs, due to the intrinsic conductive properties of gold behaving as an active layer. The authors concluded that the optimized conditions for the growth of high-density NWs were copper films of 800 nm thickness, and annealing at 450 °C for about 6 hours. The use of a conductive film on top of the CuO film performs slightly better by yielding a higher density of NWs at the lower temperature of 400 °C for about the same treatment time of 6 h.

A complex study revealing the role of gold decoration on the growth of copper oxide nanowires was carried out by Mishra *et al.*¹³⁸ The authors used the following technology sequence (Fig. 16): first, an initial array of CuO nanowires was grown until saturation using a microwave plasma at atmospheric pressure; then, the array was covered with separated gold nanoparticles; next, a second plasma oxidation treatment was carried out. It was predicted and verified that the growth of the nanowires re-started during the second oxidation. A combination of growth in plasma, where the nanowires grow quickly to the saturated length, and then the deposition of metals followed by a second exposure to plasma results in changing the morphology of the nanowires, thus providing an effective tool for control over their physical and chemical properties.

The saturation was explained using a model that took into account the mechanism of ion bombardment under the action of low-pressure plasma.¹³⁹ In contrast, the atmospheric oxygen plasma applied in the present experiment, contains a large number of radicals, electrons, dissociated oxygen atoms and

oxygen molecules; the absence of ions means that this plasma can be considered as 'mild' with respect to the damage caused by an ion-containing low-pressure plasma. The presence of the oxygen radicals results in much a more intensive oxidation of the copper sample surface, as was demonstrated in the experiments; the excited molecules cause oxidation of the copper samples at relatively low temperatures. The oxide layer usually consists of two sub-layers, namely copper(II) oxide (or cupric oxide (CuO)), and copper(I) oxide (or cuprous oxide (Cu₂O)). Both oxides are formed because of the diffusion of the species, which can be *via* lattice, boundary, or surface diffusion. In this case, the copper atoms moving from the copper layer are involved in the oxidation process on the side surface of the growing nanowire. The prolonged growth results in a decrease of the copper flow delivered from the NW base to the NW top, and the longer the NW, the lower is the supply of the copper atoms. At a certain stage the growth is terminated because all the copper is oxidized on the path to the NW top (Fig. 16). However, the intermediate deposition of noble metal enables the NW growth to be prolonged through the following mechanism. Noble metals do not interact with oxygen, thus the surface of the noble metal deposited on the side surface of the nanowires is relatively free of oxygen. More copper atoms can reach the top because the noble metal deposition decreases the density of adsorbed oxygen on the side surface of the NWs and the generation of copper oxide. As a result, copper is delivered further to the NW tip, where it is involved in a reaction with the oxygen adsorbed on the NW tip, and the NW continues to grow to a longer length.

➤ Thus, many different techniques and methods have been developed and enhanced recently to produce various types of

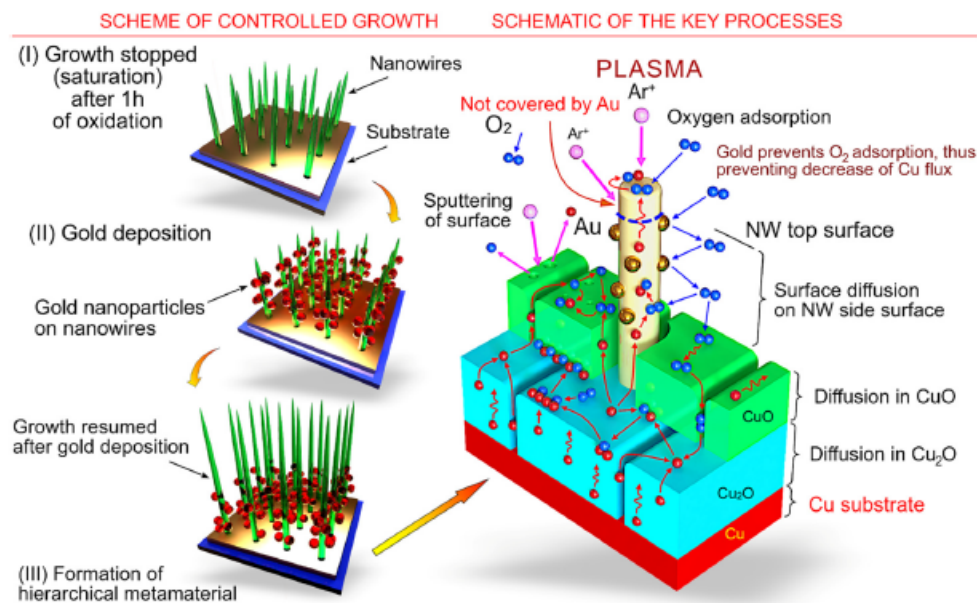


Fig. 16 (Left panel) Stages of the designed deposition-controlled growth of nanowires: (I) oxidation of copper; (II) deposition of Au nanoparticles; and (III) second oxidation. (Right panel) Formation of a hierarchical metamaterial from nanowires decorated with Au metal nanoparticles.

CuO nanostructures with low dimensionality. Below we will outline some recent innovations in the application of such structures, focusing mainly on energy, photocatalysis and sensing as the most important fields.

3. Innovations in the application of CuO nanowire-based metamaterials

First, we stress that the application of CuO nanostructures for energy, photocatalysis and sensing usually requires sophisticated functionalization to impart additional properties and ensure the high performance of CuO-based devices. The functionalization of CuO nanostructures with noble metal nanoparticles is one of the more promising techniques for enhancing the properties of CuO-based devices. Gold nanoparticles, among others, play a particularly significant role because of their catalytic activity and broad range of potential applications.¹⁴⁰ Au-based materials have many potential applications in catalysis, in oil hydro refining, as drug carriers and electrode materials, in solar cells, organic synthesis, and in water and air purification.¹⁴¹ Metal oxide-Au and graphene-Au sensors,¹⁴² photocatalytic materials,¹⁴³ photoelectrodes,¹⁴⁴ label-free biosensors,¹⁴⁵ materials for photoelectrochemical applications,¹⁴⁶ integrated electrochemical sensors,¹⁴⁷ and free-standing biosensors for the non-enzymatic detection of H_2O_2 ¹⁴⁸ are recently reported examples of metal-oxide and graphene-based metal-functionalized nanomaterials. Copper oxide nanostructure are no exception, and can often undergo similar treatment with successful practical implementations.

3.1. Innovations in CuO nanowire-based metamaterials for sensing

In the study conducted by Yang *et al.*,¹⁴⁹ sensitive CuO/Ag substrates were synthesized using thermal oxidation and magnetron-assisted sputtering. Their surface-enhanced Raman scattering (SERS)¹⁵⁰ activity was enhanced by exploiting the rich CuO NWs built on the surface of the nanostructures. Since the Ag work function is 4.1 eV and that of CuO is 5.3 eV, electrons will move to CuO when CuO NWs are covered by an Ag film, thus bringing the Fermi level to equilibrium. Therefore, one expects that combining “positively charged” Ag with “negatively charged” CuO will create a stronger electromagnetic field in response to the specific laser irradiation. Since the CuO/Ag NW array is rather sparse, the molecules prompted for the surface analysis will diffuse across the system, thus enhancing the Raman signal. It should be noted that in the future, copper oxide nanostructures may be used even more efficiently for signal amplification in the SERS-based detection of biomolecules with Au and other noble metals films through hot-spot field enhancement.¹⁵¹

CuO nanoneedles on a commercial ceramic tube¹¹³ were tested as an excellent H_2S gas sensor. The authors concluded that further improvement could be made by designing heterojunctions supplemented with suitable modification of CuO the NW arrays (Fig. 17). Hierarchical nanostructures built upon Au nanoparticles and CuO NWs have been implemented by

RECENT INNOVATIONS IN CuO NANOSTRUCTURE APPLICATIONS: GAS SENSING

CuO nanoneedle array based sensor in air and in H_2S

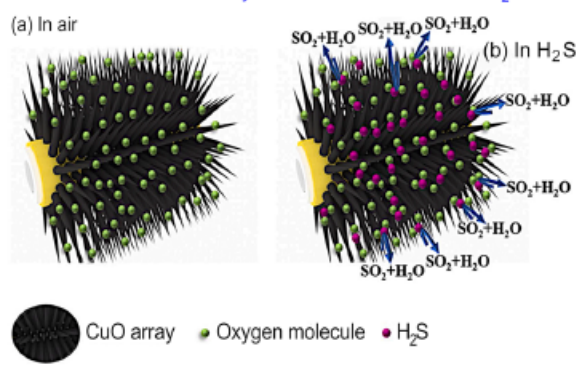


Fig. 17 CuO nanoneedle array-based H_2S gas sensor. (a) Sensor based on CuO nanoneedle arrays in air, and (b) in H_2S . In addition, the system performance can be further improved by building a heterojunction, followed by suitable noble metal modification of the CuO nanowire arrays. Reprinted with permission from Hu *et al.*, ref. 113. Copyright 2021, Elsevier.

Zhao *et al.*¹⁵² for use as portable enzymeless glucose detectors. Le *et al.*¹²⁸ successfully developed such hybrid systems as biosensors, aimed at α -cysteine detection, displaying properties of a wide detection range, lasting stability, and excellent selectivity. The emerging synergistic effects boosted the sensing performance by improving the exposure of the relevant electroactive sites, accelerating the rate of charge transfer, and increasing the specific surface area. The latter was the result of a prominent 3D hierarchical nanoarchitecture, characterized by impressive mechanical properties that enhance the electrolyte penetration, the diffusion of ions, mass transfer, and the ability to increase charge transfer.

Aluminum-doped CuO nanoleaves can also be used for the development of carbon monoxide sensors (Fig. 18)¹⁵³ A coprecipitation technology was employed to produce doped copper oxide nanostructures, which were then employed as basic building blocks for the fabrication of cheap sensors for the detection of carbon monoxide with high sensitivity at low working temperatures. The sensitivity of the CO sensors to various gas concentrations was investigated over a very wide range of 35 to 12 500 ppm, and at temperatures in the range of 70–230 °C. While pure CuO nanostructures showed a response of about 50% at 120 °C, the response of aluminum-doped nanoleaf-shaped copper oxide structures was substantially stronger, at nearly 70% at 150 °C.

Finally, Hao *et al.* modified Cu NWs by depositing them on top of graphene oxide nanosheets to obtain a sensor for multiple simultaneous uses, such as in the detection of ascorbic acid, dopamine, and acetaminophen.¹⁵⁵ Fig. 19a presents a schematic of analyte gas molecule adsorption on the CuO/rGO hybrid surface. The novel copper oxide/functionalized graphene hybrid nanostructures were fabricated for gas sensing.¹⁵⁴ Fig. 19b shows the sensing response curve of a quartz crystal microbalance gas sensor with the CuO/rGO hybrid.

RECENT INNOVATIONS IN CuO NANOSTRUCTURE APPLICATIONS

CuO nanoleaves for carbon monoxide gas sensing

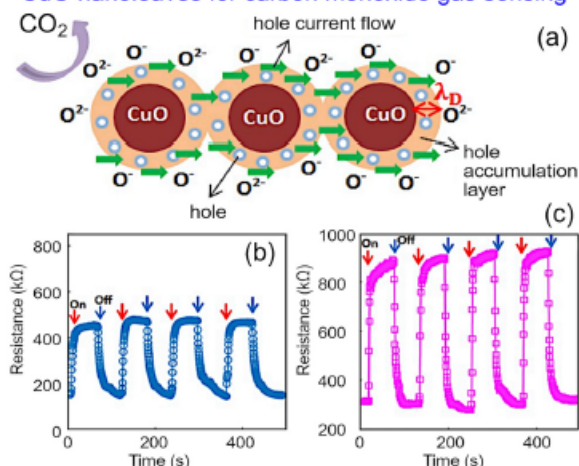


Fig. 18 Carbon monoxide sensor. (a) Principle of the sensing mechanism of copper oxide for carbon monoxide utilizing a hole accumulation layer. (b and c) Repeated response to 800 ppm carbon monoxide for (b) pure CuO sensors; and (c) Al doped CuO sensors when operated at their optimum temperature of 120 and 150 °C, respectively. Reprinted with permission from Molavi *et al.*, ref. 153. Copyright 2020, Elsevier.

3.2. Innovations in CuO nanostructure-based metamaterials for biosensing

Biosensing is a very important area of application where CuO-based nanostructures and metamaterials ensure excellent results. CuO-based nanostructures were used recently for the fabrication of quite different biosensors¹⁵⁶ that are capable of detecting neuromodulatory molecules such as dopamine,^{157,158} as well as active substances that are important for living beings, *e.g.*, glucose, cysteine, carbon monoxide, hydrogen peroxide

and others. Below we provide a brief overview of some of the recent innovations in copper oxide-based biosensor technology.

A colorimetric visual biosensor for detecting L-cysteine has recently been synthesized using dopamine-coated copper oxide nanoparticles. This innovative approach is based on the use of natural dopamine, an organic molecule from the catecholamine and phenethylamine families and a neuromodulator, the chemistry of which is frequently used for the surface functionalization of copper oxide particles to improve their stability and sensitivity to L-cysteine (Fig. 20a).¹⁵⁸ As a semi-essential amino acid that plays an important role as a building block in protein synthesis, L-cysteine occurs at intracellular concentrations of 20–200 μM in biosystems.¹⁵⁹ The dopamine-coated copper oxide nanoparticles (CuO@DOP NPs) were fabricated using microwave radiation (operating parameters: 800 W, 2450 MHz). The actual suitability of the newly developed sensor for the detection of L-cysteine was ascertained by using it to analyse clinical samples of blood serum and urine. Fig. 20b shows the representative fluorescence emission spectra of the dopamine-coated copper oxide nanoparticles as a function of the L-cysteine concentration. The authors conclude that a sensor based on such dopamine-coated copper oxide nanoparticles may be used for the detection of L-cysteine both rapidly and with a high level of sensitivity across a broad range of concentrations, while being biocompatible with living systems.

Nanostructures with interesting and very promising properties were recently reported by Cheng *et al.*¹⁶⁰ Mesoporous hollow CuO spheres were fabricated and tested for use in paper-based hydrogen peroxide sensors, where the research was stimulated by the increasing demand for reliable, convenient and cheap paper-based sensors for the rapid detection of hydrogen peroxide in clinical, personal and industrial settings. Fig. 21a illustrates the schematic for the fabrication of hollow mesoporous CuO sphere nanozymes. The latter were fabricated by decomposing spherical copper-polyphenol colloidal particles. The resulting spherical structures were uniform in size (at ~100 nm) and hollow, and

RECENT INNOVATIONS IN CuO NANOSTRUCTURE APPLICATIONS: SENSORS

Copper oxide / functionalized graphene hybrid for gas sensing

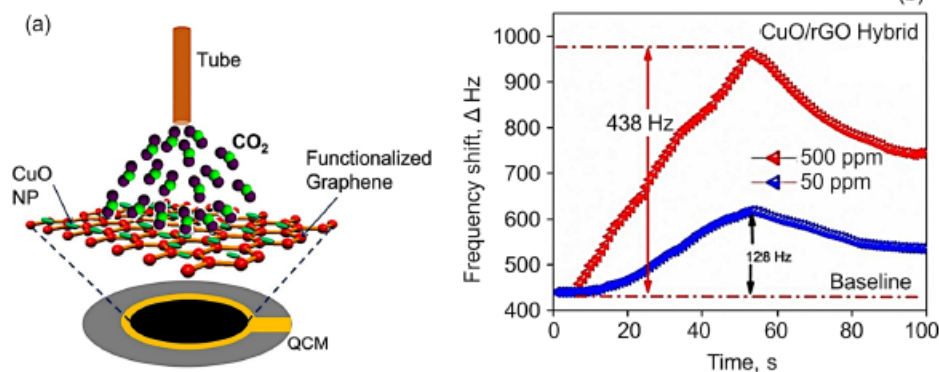


Fig. 19 Copper oxide/functionalized graphene hybrid for CO₂ gas sensing. (a) Schematic of analyte gas molecule adsorption on the surface. (b) Sensing response curve of the quartz crystal microbalance. Reprinted from Gupta *et al.* 2022, ref. 154 under terms and conditions of CC BY license.

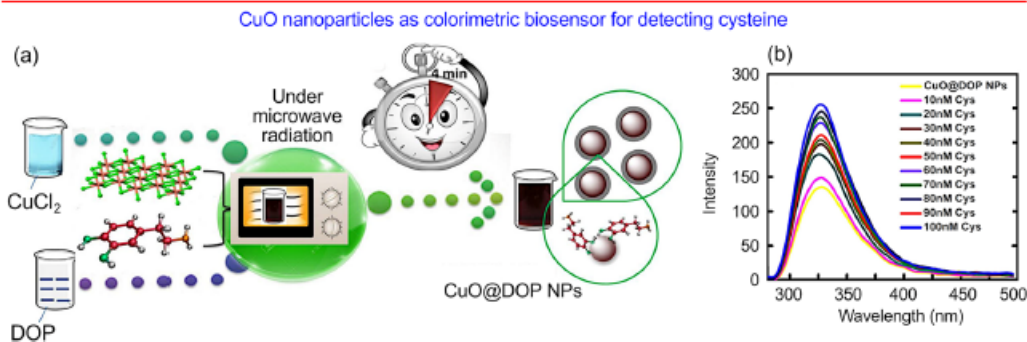


Fig. 20 CuO nanomaterial-based biosensor. (a) CuO nanoparticles as a colorimetric biosensor for detecting cysteine, and (b) fluorescence spectra as a function of the wavelength for dopamine-coated copper oxide nanoparticles, for different concentrations of L-cysteine. Reprinted with permission from Rohilla *et al.*, ref. 158. Copyright 2020, Elsevier.

featured an impressive specific surface area that reaching about $58 \text{ m}^2 \text{ g}^{-1}$, with significant (about $0.5 \text{ cm}^3 \text{ g}^{-1}$) porosity, and interconnected mesopores reaching diameters of 5.8 nm.

Fig. 21b shows a schematic of the detection mechanism for hydrogen peroxide by the hollow mesoporous CuO sphere

nanozymes. The sensors demonstrated an excellent peroxidase-detection activity as well as detection repeatability, as illustrated in Fig. 21c. They can be used for the colorimetric detection of peroxidase in the concentration range of 2.4–150 μM .

A method for the fabrication of hydrogen peroxide sensors using a simple polyol method, where copper oxide particles are produced when copper(II) acetate is hydrolyzed in 1,4-butanediol in the absence of additives, was also reported recently.¹⁶¹ Fig. 22 illustrates the general approach for the synthesis of the copper oxide nanoparticle (CuONP)-based non-enzymatic electrochemical sensor. The authors simply drop-cast the sensing CuONP nanomaterial onto a glassy carbon electrode (GCE) which was modified using the poly(3,4-ethylenedioxythiophene) (PEDOT) conducting polymer. The latter was formed as a layer on a glassy carbon electrode *via* a sinusoidal voltage method.

The formation of nanoparticles assembled into quasi-prismatically-shaped microscale aggregate structures with sizes in the range from 1 μm to $\sim 200 \mu\text{m}$, characterized by a mesoporous structure, was reported recently. Furthermore, the newly designed electrochemical sensing platforms appear to respond linearly with an increase in the hydrogen peroxide concentration from 0.04 to 10 mM, and have a low threshold (about 8.5 μM) for peroxide detection. Importantly, the sensing platform features very good anti-interference characteristics when substances, *e.g.*, potassium nitrate and potassium nitrite, which are known to interfere with the detection of hydrogen peroxide, are present.

In addition to detecting biomolecules, copper oxide-based biosensors may be useful in the detection of viral protein structures for both health and environmental applications.¹⁶² Viral infections can spread rapidly amongst human, animal and plant populations, and their timely detection is often critical to limit their spread and develop a treatment plan. Amongst the most prominent examples of this are the human immunodeficiency virus (HIV), the Ebola virus, and, most recently, COVID-19. The polymerase chain reaction (PCR) is currently the most reliable technology used to determine the identity of a virus; however, its use is limited by its high cost and the length of time it takes to produce the answer, especially when demand for the service

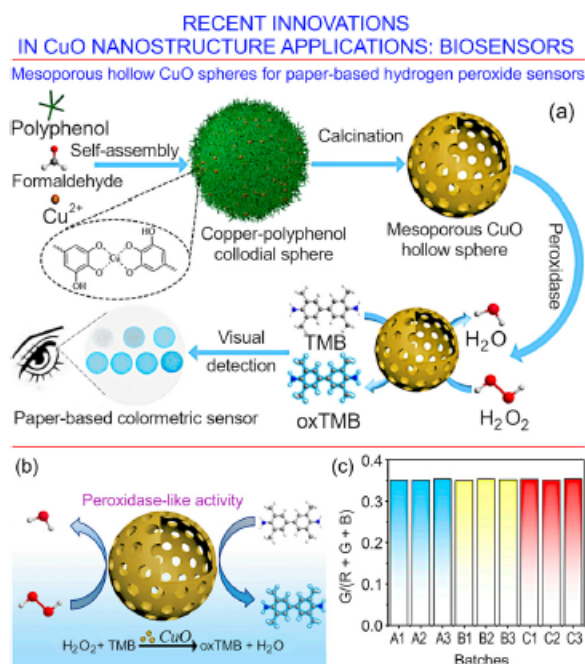


Fig. 21 Mesoporous hollow CuO spheres for paper-based hydrogen peroxide sensors. (a) Schematic of the fabrication of hollow mesoporous CuO sphere nanozymes, and the application of this material as a paper-based sensor for the colorimetric detection of hydrogen peroxide. (b) Schematic of the detection mechanism for hydrogen peroxide by the hollow mesoporous CuO sphere nanozymes. (c) Measurement repeatability for three batches of the paper-based sensors towards the detection of hydrogen peroxide. Reprinted with permission from Cheng *et al.* 2021, ref. 160 under terms and conditions of CC BY license.

RECENT INNOVATIONS IN CuO NANOSTRUCTURE APPLICATIONS: BIOSENSORS

Copper oxide nanoparticles for hydrogen peroxide sensing

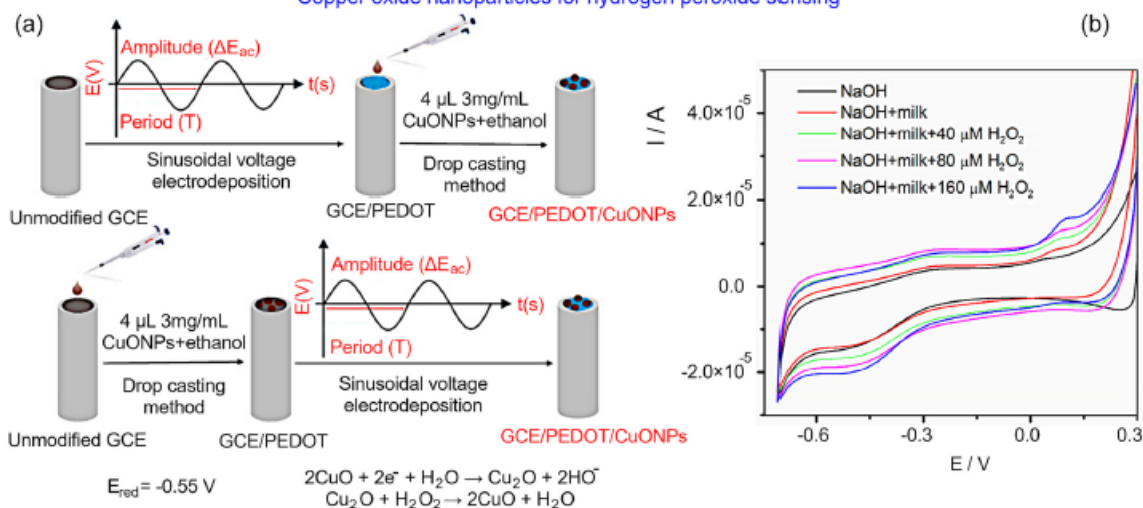


Fig. 22 Copper oxides for biosensing. (a) Approach for the synthesis of a copper oxide nanoparticle (CuONP)-based non-enzymatic electrochemical sensor for the detection of hydrogen peroxide. (b) Cyclic voltammetry results for the sensor based on GCE/PEDOT-CuONPs when used to detect different amounts of hydrogen peroxide in a diluted milk sample. Reprinted with permission from Lete et al. 2022, ref. 161 under terms and conditions of CC BY license.

is high. To circumvent these limitations, a new generation of sensors is needed. An example of such a sensor was recently developed using a graphite electrode decorated with CuO nanostructures deposited *via* wet chemical processing. The antibody corresponding to the nucleocapsid protein of the SARS-CoV-2 virus was immobilized on the CuO-graphite electrode structure, where the N protein is an abundant RNA-binding protein that plays an

important role in genome packaging in viruses. The antigen-antibody interaction was confirmed using square wave voltammetry (SWV). Fig. 23a illustrates the fabrication technology and explains the main steps, and Fig. 23b shows stronger performance brought about when graphite electrodes are modified with both hydroxyapatite (HAP) and copper oxide (CuO), compared with a pure graphite electrode or that modified with HAP alone.

RECENT INNOVATIONS IN CuO NANOSTRUCTURE APPLICATIONS: BIOSENSORS

Copper oxide modified electrodes for biosensors for determination of environmental viral structures

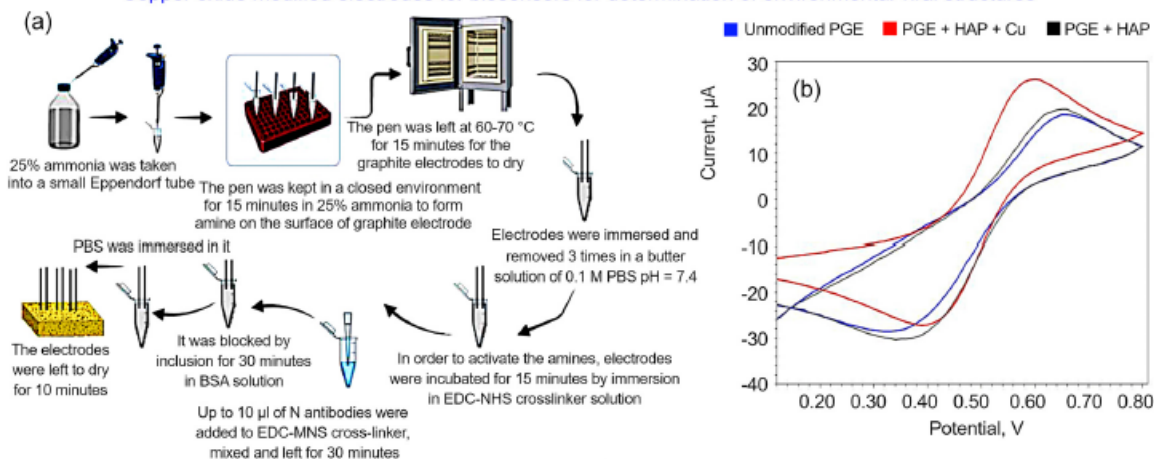


Fig. 23 Copper oxides for biosensing. (a) Functionalization of pen shaped graphite electrodes endowed with hydroxyapatite and copper oxide carrying a nucleocapsid of COVID-19 antibody, and an illustration of the main pathways in the preparation of the electrochemical sensor based on copper oxide. (b) Results from cyclic voltammetry applied to untreated and treated pen shaped graphite electrodes with hydroxyapatite and copper oxide for a solution of 2.0 mM FeCl₃ in 1.0 M HCl using a 10 mV s⁻¹ scanning rate. Reprinted with permission from Elkici et al. 2022, ref. 162 under terms and conditions of CC BY license.

3.3. Innovations in CuO nanowire-based metamaterials for glucose detection

The quick and reliable detection of glucose remains a very important challenge as the incidence of diabetes is increasing at an unprecedented rate all over the globe. At present, close to half a billion people on the planet suffer from this chronic condition, as estimated by the World Health Organization, while only about one hundred million cases were registered in 1980. Our ability to deal with this disease relies on the availability of devices and technologies for the accurate, reproducible and rapid detection of glucose concentrations of in a variety of media, from patient blood to food and medication that the patient may be consuming. Unfortunately, the currently available sensors for glucose detection are limited by their use of high-cost enzymes, and the results that they produce are insufficiently reproducible. As a consequence, efforts have been made to develop cheaper non-enzymatic sensors with both high selectivity and sensitivity.¹⁶³

Baek *et al.*¹⁶⁴ proposed a novel electrospinning method for the fabrication of a graphene-based oxide nanofiber decorated with Cu-nanoflowers to be employed as an electrochemical biosensor for glucose detection (Fig. 24). Chen *et al.*¹⁶⁵ fabricated an NO₂ sensor using gold-functionalized CuO nanorods *via* a hydrothermal process at a room temperature. The resulting CuO nanorods had a diameter of 30–60 nm and a length of 150–200 nm, and displayed a robust structure that was not affected by the Au functionalization process used. It was found that small Au nanoparticles, with a diameter smaller than about 20 nm, were quite uniformly distributed over the surface. The authors concluded that the observed enhanced sensing

characteristics were due the increased number of oxygen vacancies as well as to the lower activation energy as a result of Au functionalization. Lee *et al.*¹⁶⁶ developed a network of p-CuO NWs grown on a predefined electrode pad pattern *via* the thermal oxidation of a Cu layer deposited using an electron-beam. Heat treatment of the Au layer, deposited by sputtering, was used to functionalize the Au nanoparticles. The latter had a diameter in the range of 20–145 nm, obtained by controlling the deposited Au layers thickness. It was found that Au functionalization enhanced the sensitivity of the CuO nanowires to a large extent, yielding the best sensing characteristics after functionalization with 60 nm-diameter Au NPs. One may conclude that, for functionalization to be efficient, the size optimization of the nanoparticles is very important to ensure the superior sensing characteristics of the oxide nanowires. Indeed, the response for CO was improved by approximately a factor of two compared with the traditional method using benzene and toluene.

Nanowires with high aspect ratios were reported by Hadiyan *et al.*¹⁶⁷ Cu₂O NWs of 180 nm in diameter and 8 μm in length were synthesized *via* electrodeposition onto a polycarbonate substrate immersed in an aqueous solution. Further annealing yielded CuO/Cu₂O composite NWs. They found that, at 250 °C, the Cu₂O NWs work well as an accurate ethanol gas sensor, while their CuO/Cu₂O counterparts also showed good sensing response for CO at 150 °C. These results indicate that the Cu₂O NW gas sensor is almost six times faster than sensors designed using CuO/Cu₂O nanowires. A comprehensive analysis conducted by Qi *et al.*¹⁶⁸ on CO oxidation both in oxygen-rich and in hydrogen-rich atmospheres demonstrated that the CuO species play a major role in the oxidation of CO, consistent with the fact that the superior Au/CuO catalyst relies on its strong interaction with the substrate gold nanoparticles.

An excess glucose concentration in human blood can have a negative impact on human health. Therefore, people are actively working on the issue of detecting blood glucose contaminants in the human body, which can be an important indicator of the surge of diabetes. A complex copper oxide/zinc oxide nanostructured material for non-enzymatic glucose sensing was reported recently by Cheng *et al.*¹⁷¹ Electrodes were fabricated *via* a simple solution method, *i.e.*, by the immersion of ZnO nanostructures into copper ion solutions. Tetrapod-like ZnO nano-powders were first obtained using a direct current plasma method, and the resulting surface was uniformly decorated with CuO nanocrystalline particles (Fig. 25a). Fig. 25b and c illustrate the response of the fabricated sensor to glucose.

In the interesting work by Patil *et al.*, thin CuO films synthesized *via* chemical bath technology were reported.¹⁶⁹ They found that thin films feature a much stronger response to glucose, in comparison with their response to ascorbic acid. The work by Steinhauer reviewed the recent achievements in designing novel devices using copper oxide nanomaterials, and discussed the key factors that determine the sensing characteristics and efficiency.¹⁷⁰

Recently, an interesting electrodeposition approach was demonstrated to design ultrasensitive amperometric non-enzymatic

RECENT INNOVATIONS IN CuO NANOSTRUCTURE APPLICATIONS

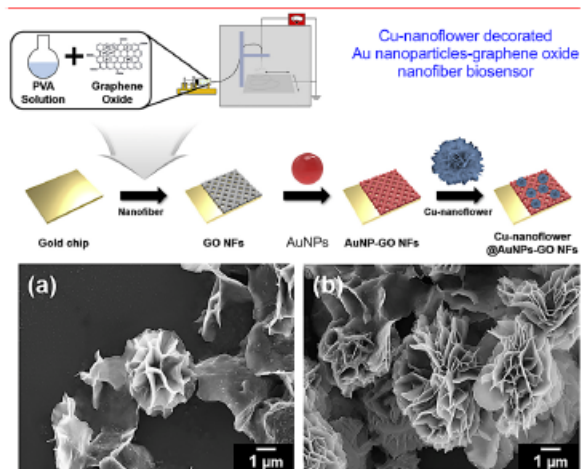


Fig. 24 Electrochemical biosensor for glucose detection. (Upper panel) Synthesis of copper (Cu)-nanoflowers, decorated on graphene oxide (GO) nanofibers with gold nanoparticles, as an electrochemical biosensor for the detection of glucose. (Bottom panels) Optimization of the Cu-nanoflower at different GO concentrations: (a) 0.3 mg mL⁻¹, and (b) 0.5 mg mL⁻¹. Reprinted with permission from Baek *et al.*, ref. 164. Copyright 2021, Elsevier.

RECENT INNOVATIONS IN CuO NANOSTRUCTURE APPLICATIONS: SENSORS

Copper oxide / zinc oxide nanostructures for non-enzymatic glucose sensor

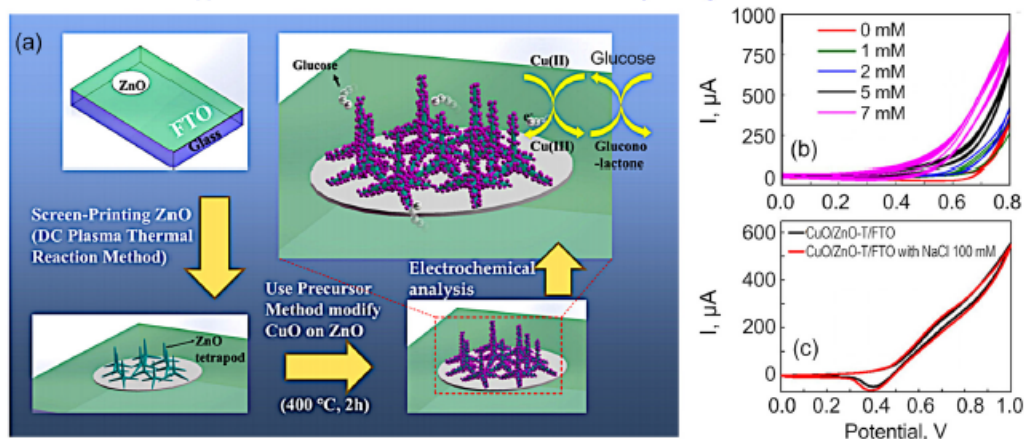


Fig. 25 Copper oxide/zinc oxide nanostructures for a non-enzymatic glucose sensor. (a) Schematic of the fabrication process. (b) Current–voltage curves for glucose at various concentrations, and (c) response of the sensor to glucose solution. Reprinted from Cheng *et al.* 2021, ref. 171 under terms and conditions of CC BY license.

glucose biosensors that use a more complex architecture, *i.e.*, magnetically active thiol-functionalized $\text{Fe}_3\text{O}_4\text{-SH}$ over a glassy carbon substrate, decorated with honeycomb-like copper oxide onto which silver nanoparticles are immobilized.¹⁷² Fig. 26a illustrates the electrode structure that incorporates all of these elements, and Fig. 26b shows the cyclic voltammograms of glucose in NaOH for the bare electrode and for various CuO-honeycomb-supported systems. The thus-fabricated sensors show excellent sensitivity for glucose and a stability of response over a broad range (0.06–1000 mM) of glucose concentrations, with a detection threshold of 15 nM. As such, these sensors could potentially be used to detect glucose reliably in blood and urine clinical samples.

One more technology for the fabrication of copper oxide-based biosensors for non-invasive glucose detection was developed recently by Franco *et al.*¹⁷³ In order to fabricate the sensor, the authors first print-coated electrodes onto fabric that was made of cellulose using a paste containing graphene. Cellulose is an excellent substrate material as it offers an attractive combination of minimal cost and toxicity, abundance, excellent compatibility with living systems, the ability to be disposed of and degrade naturally, as well as excellent porosity that maximises the contact area between the sensor and the biofluid. Drop-casting was then carried out to deposit Cu_2O nanoclusters onto the surfaces of the graphene electrodes, with the Cu_2O crystals synthesized *via* an ascorbic acid reduction method. The sensing was made possible by exploiting electron transfer from copper oxide in an aqueous solution of sodium hydroxide and artificial sweat that takes place at a considerably low potential of +0.35 V, with the existence of the reduction–oxidation reaction between glucose and the nanoparticles confirmed using cyclic voltammetry and other characterization methods.

Fig. 27a illustrates the simplicity of this fabrication technology, which makes it highly attractive for the large-scale production of

cheap disposable sensors that at the same time demonstrate a high level of sensitivity and reproducibility. The latter characteristics are shown in Fig. 27b for a NaOH solution with glucose

RECENT INNOVATIONS IN CuO NANOSTRUCTURE APPLICATIONS: GLUCOSE SENSORS

Nanocomposite films of CuO honeycombs - Ag nanoparticles for glucose sensing

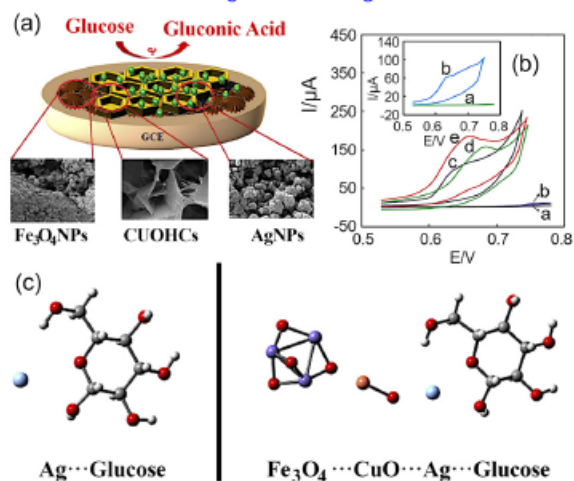


Fig. 26 CuO honeycombs and nanoleaves for glucose and carbon monoxide sensing. (a) Electrode fabricated on nanocomposites of CuO honeycombs/Ag nanoparticles. (b) Cyclic voltammograms, for 0.1 mM glucose in 0.1 M NaOH solution, on various electrodes, including a bare electrode and various CuO-honeycomb-supported systems (see the details about traces labelled a–e in the original publication). (c) Optimized structures of the complexes: $\text{Ag} \cdots \text{glucose}$ and $\text{Fe}_3\text{O}_4 \cdots \text{CuO} \cdots \text{Ag} \cdots \text{glucose}$. Reprinted with permission from Baghayeri *et al.*, ref. 172. Copyright 2020, Elsevier.

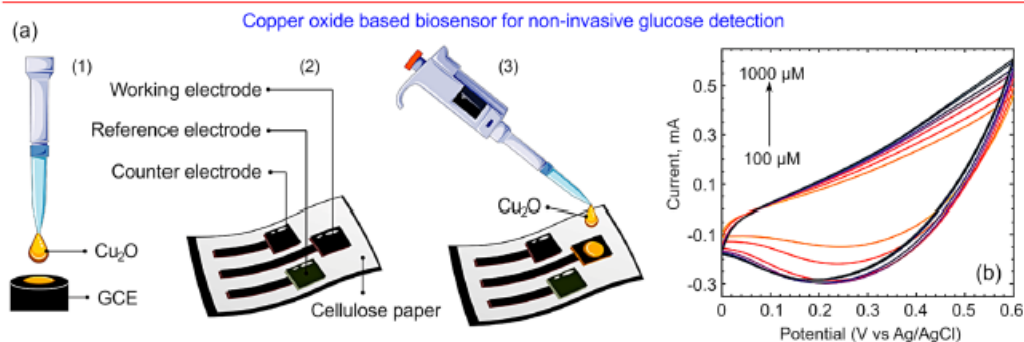


Fig. 27 Copper oxide-based biosensor for non-invasive glucose detection. (a) General approach for the preparation of glucose-sensing platforms on a glassy carbon electrode (1; used for material characterization) and on a graphene electrode on a flexible glucose substrate (2–3; as a proof-of-concept study). Ag/AgCl is used as a pseudo-reference electrode. (b) Cyclic voltammetry curves for a sensor based on the Cu_2O nanocluster-decorated graphene working electrode shown in (a3) for a glucose concentration in the range of 100 to 1000 μM in 0.1 M NaOH solution. Reprinted with permission from Franco *et al.* 2022, ref. 173 under terms and conditions of CC BY license.

levels ranging from 0.1 to 1 mM. Under these conditions, sensors based on copper oxide over a glassy carbon electrode show a sensitivity of about $1082 \mu\text{A mM}^{-1} \text{cm}^{-2}$, whereas those based on copper oxide nanoclusters over graphene electrodes (on cellulose) recorded a sensitivity of about $182 \mu\text{A mM}^{-1} \text{cm}^{-2}$. These measurements confirm the potential for these sensors to find applications in the non-invasive monitoring of patients with blood sugar disorders, particularly where portability improves patient uptake and compliance, as they show selectivity towards glucose in the presence of interfering urea and NaCl that are typically present in human sweat.

3.4. Innovations in CuO nanowire-based metamaterials for energy and catalysis

Many experiments have been published that report the use of gold as a metal nanoparticle source for the functionalization of CuO nanostructures. Efficient CuO photocathodes that feature remarkable stability were reported by Masudy-Panah *et al.*¹⁷⁴ These authors exploited the novel engineering of *in situ* materials based on the deposition of gold-palladium (Au-Pd) nanoparticles on the surface of CuO. The basic functioning of these nanostructures relies on plasmonic centers which trap the incoming light, thus enhancing the optical absorption of the system. The associated hydrogen amount becomes larger than in the cases of bare CuO photocathodes.

The results obtained by Sun *et al.* suggest that complex materials of hybridized cuprous sulfide and copper may play an important role in water splitting in pre-catalytic processes that do not use noble metals.⁸⁵ Interestingly, the Cu_2S particles evolved into CuO nanowires directly in the oxygen evolution reaction (OER). Similarly, nanostructures for the OER were reported by Butt *et al.*, and the copper nanowires were fabricated *via* the reduction of CuO nanowires in hydrogen plasma.¹⁷⁵ Testing this material for application in the hydrogen evolution reaction (HER) proved its exceptional characteristics, with the performance significantly exceeding the values obtained for the bulk copper material. Next, the complex

$\text{CuO/Cu}_2\text{O}$ nanomaterial for Na^+ ion storage was synthesized by Ma *et al.*¹⁷⁶ *via* the *in situ* modification of copper foam using the hydrothermal method (Fig. 28). The synergistic effect of CuO and Cu_2O oxides can contribute to decreasing the volume change between the host materials, and, as a result, additional Na^+ ions can be stored in the interfaces between the two oxides, leading to conspicuous capacity enhancements. The review by Majumdar *et al.*¹⁷⁷ outlines the recent advancements in CuO-based nanosystems, which are typically employed for the design of better electrode materials aimed at developing advanced supercapacitors. In addition, they discuss the electrochemical responses for a variety of CuO nanostructures.

Pristine nanomaterials based on copper oxides feature a relatively low electrical conductivity and a rather low electrochemical stability. To overcome these and some other shortcomings, CuO nanomaterials could be combined with pseudo-capacitive materials such as, *e.g.*, chalcogenide or carbon-containing materials like conductive polymers, graphene- and carbon nanotube-containing materials. A combination of CuO nanowires with graphene sheets was recently demonstrated to obtain highly porous, conductive hybrid nanomaterials that could be fruitfully employed as high-performance supercapacitor electrodes with enhanced capacitances (Fig. 29).¹⁷⁸ Currently, the development of new technology based on high-performance nanoelectromechanical systems (NEMS) is limited by the problems associated with the synthesis of NEM switches based on single nanowires. A successful application of CuO nanowires as a NEM switch element was recently reported by Jasulaneca *et al.*¹⁷⁹ Moreover, the anodized copper foams obtained by Abd-Elnaem *et al.* have been to be an efficient material for enhancing the efficiency of organic dye decomposition.¹⁸⁰

A number of experiments carried out to develop a method for the highly-controllable synthesis of nanostructures includes a stage achieves metal nanoparticle deposition. In the study reported by Kim *et al.*, broadband absorbing surfaces for solar energy applications were developed on self-aggregated alumina nanowires.¹⁸¹ Excellent antibacterial activity was reported by

RECENT INNOVATIONS IN CuO NANOSTRUCTURE APPLICATIONS: BATTERIES

Porous copper oxide heterostructured arrays for high-performance batteries

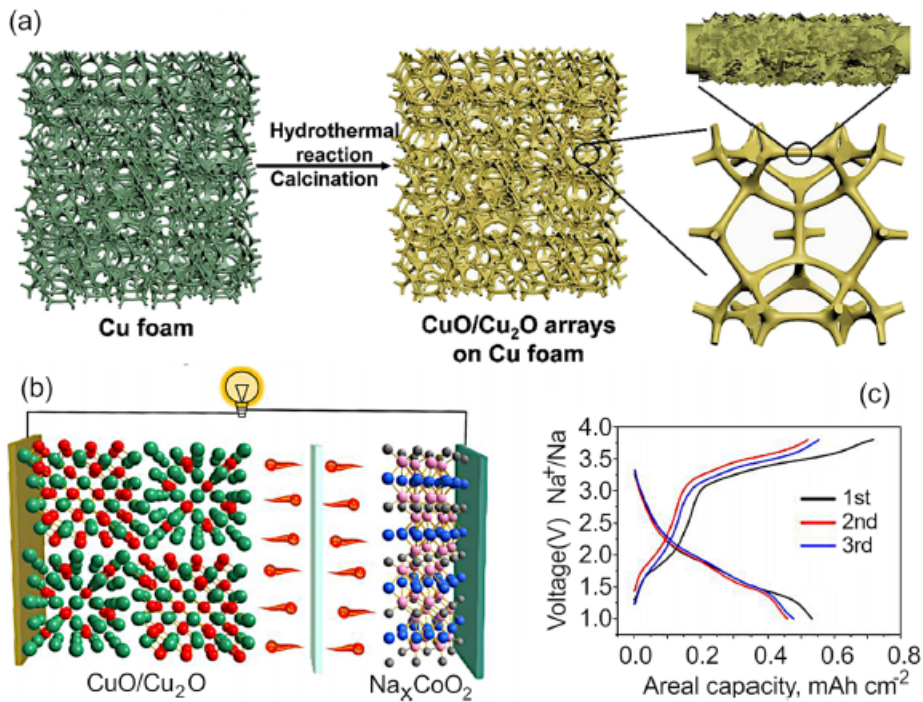


Fig. 28 Porous copper oxide heterostructured arrays for high-performance batteries. (a) Synthesis of CuO/Cu₂O arrays. (b) Full sodium-ion battery with a CuO/Cu₂O anode and an Na_xCoO₂ cathode. (c) Initial three charge–discharge curves for a (CuO/Cu₂O)–(Na_xCoO₂) full battery operating at a current density of 0.5 mA cm⁻². Reprinted with permission from Ma *et al.*, ref. 176. Copyright 2021, Springer.

Menazea *et al.* for the material based on graphene oxides with the gold and CuO nanoparticles.¹⁸² Sun *et al.* proved that the functionalization of transition metal surfaces, carrying dichalcogenide nanosheets, could be used to obtain better electrical contacts to devices, together with the improvement of their catalytic and sensing characteristics.¹⁸³ Along this line of research, Attar *et al.*¹⁸⁴ reported, for the first time, the high

electrocatalytic performance of a carbon paste electrode modified by polypyrrole–copper oxide particles (Cu₂O/PPy/CPE) for the electrocatalytic oxidation of ethanol.

Fig. 30a shows the technology for the fabrication of self-supporting electrodes for the oxygen evolution reaction and supercapacitors. By employing a facile electro-oxidation method, Cu_xO nanoflakes have been successfully grown *in situ* on a

RECENT INNOVATIONS IN CuO NANOWIRE APPLICATIONS

Copper oxide nanowire/graphene hybrid nanostructure for high-performance supercapacitors

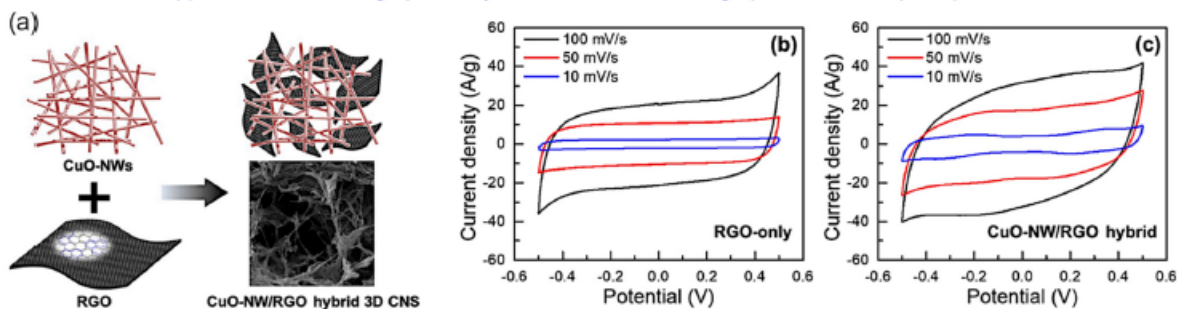


Fig. 29 CuO-nanowire/reduced graphene oxide hybrid for high-performance supercapacitors. (a) Schematic of the CuO-nanowire/reduced graphene oxide hybrid three-dimensional carbon nanostructures. (b) and (c) Electrochemical current–voltage curves for RGO-only and CuO-nanowire/reduced graphene oxide hybrid three-dimensional carbon nanostructures. Reprinted with permission from Luan *et al.*, ref. 178. Copyright 2019, Elsevier.

Copper oxide nanowire-based nanostructures for high-performance supercapacitors

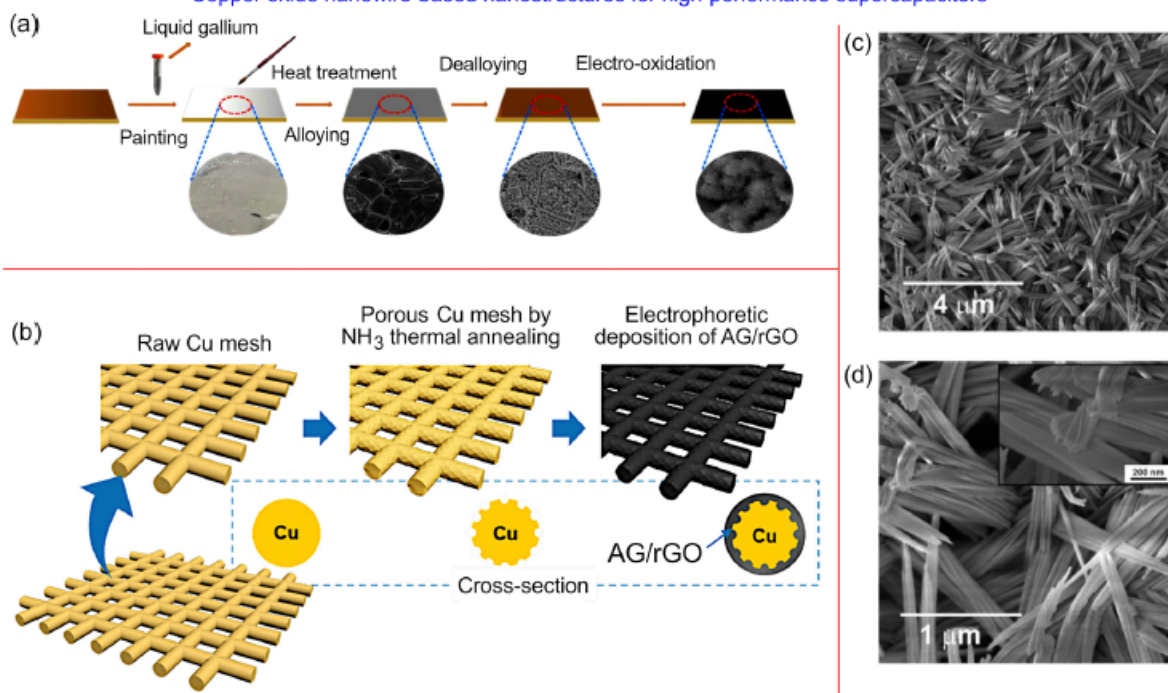


Fig. 30 (a) Fabrication process of the Cu_xO/Cu electrodes. Reprinted from Li *et al.* 2022, ref. 185 under the terms and conditions of the CC BY license. (b) Fabrication of activated graphene (AG)/reduced graphene oxide (rGO) deposited on a porous Cu (P-Cu) mesh. Reprinted from Lim *et al.* 2021, ref. 186 under the terms and conditions of the CC BY license. (c and d) Morphology of anodic CuO/Cu₂O nanoneedles obtained for Cu foils anodized in KOH-H₂O-NH₄F electrolyte. Inset in (d) is a high-magnification SEM image. Reprinted from Oyarzún *et al.* 2021, ref. 187 Copyright under the terms and conditions of the CC BY license.

nanoporous Cu foil. In order to realize the formation of a nanoporous Cu layer on a flexible Cu foil, a Ga-Cu alloying/dealloying system prepared on a Ga-assisted surface was used. Electro-oxidation carried out at a constant potential was used to modify the nanoporous Cu layer containing the Cu_xO nanoflakes. The optimum electrode configuration was able to deliver a rather high areal capacitance of about 0.75 F cm⁻² at a current density of 0.2 mA cm⁻². The samples were able to maintain about 95% of the capacitance through 12 000 cycles. The supercapacitor built using Cu_xO/Cu as the positive electrode and activated carbon as the negative electrode demonstrated energy and power densities of 24.2 W h kg⁻¹ and 0.65 kW kg⁻¹, respectively.¹⁸⁵

Another technology for CuO-based supercapacitors has been reported recently by Lim *et al.*, in which a porous Cu mesh was implemented as the current collector, while a porous surface was made *via* thermal annealing of the Cu mesh in an ammonia gas environment.¹⁸⁶ Activated graphene, with a hierarchical porous distribution and a high specific surface area, and graphene oxide were deposited onto the porous Cu substrate *via* electrophoretic deposition. Subsequently, the graphene oxide was transformed into electrically conductive rGO. This

material yielded a high specific capacitance reaching 140 F g⁻¹, together with a prominent energy density exceeding 3 W h kg⁻¹ (Fig. 30b). Fig. 30c and d show SEM images of the anodic CuO/Cu₂O nanoneedles obtained for Cu foils anodized in KOH-H₂O-NH₄F electrolyte.¹⁸⁷

Thus, various types of CuO nanostructures with low dimensionality could be successfully implemented for advanced applications in sensing, energy and catalysis, as follows from the recent achievements (mostly 2021 and 2022) reviewed above. Apparently, this material still holds great potential for future developments, and more effort should be devoted to the design of novel, complex, hierarchical materials and metamaterials for advancing future applications.

However, apart from the above outlined sensing, energy and catalysis applications, CuO nanostructures with low dimensionality could also be implemented in other leading-edge applications, such as space exploration and medicine. In the Outlook section below we will briefly discuss these novel areas of application for CuO nanostructures.

Furthermore, some further problems and open questions related to the CuO nanostructures will be outlined in the Outlook section.

4. Outlook

Diversity of technologies and applications for CuO-based low-dimensional nanostructures

The above-reviewed recent results on the development of fabrication technologies and advanced applications of CuO-based low-dimensional nanostructures demonstrate a flourishing diversity of techniques and devices that utilize CuO-containing materials. Importantly, many types of complex, hierarchical materials systems and metamaterials could be built using CuO-based low-dimensional nanostructures. The spectrum of recently reported morphologies and applications of low-dimensional CuO nanostructures, as examined in this review, is presented in Fig. 31. To get a better overview of this diversity and thus to stress the greatest potential of CuO-based low-dimensional nanostructures for future applications, we have compiled the general list in Table 1 that provides examples that have been discussed in detail in this review.

From this table one can see that the most commonly discussed CuO nanostructures are nanowires, while some unique shapes, such as butterfly-like nanostructures, porous hollow nanotubes, nano-needle arrays, Cu_xO nanoflakes,

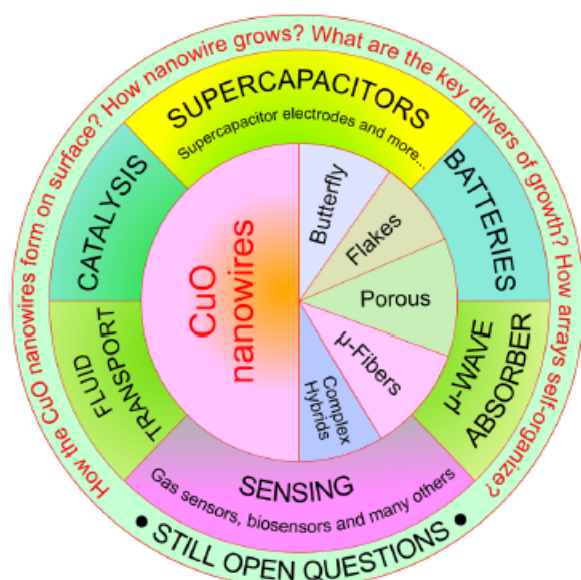


Fig. 31 Spectrum of recently reported morphologies and applications of low-dimensional CuO nanostructures examined in this review. While energy, sensing and catalysis could be considered as the 'classic triad of applications' for CuO nanomaterials, other applications have also been the focus of recent investigations. By contrast, nanowires and large organised arrays of CuO nanowires are the most common form of CuO nanomaterials for advanced applications, but other morphologies including quite exotic forms, such as butterfly-like nanostructures, have also been the subject of focus. Besides, in spite of great advances achieved in the synthesis and applications of CuO nanomaterials for advanced applications and theoretical insights in the growth mechanisms, some questions are still remain, as highlighted within this scheme, with the most intriguing queries being "What are the key drivers of growth?" and "How arrays of CuO nanowires self-organize during growth on the surface?".

porous mesh and microfibers, have also been created and successfully tested for applications in the past 2 years. Complex CuO/graphene and CuO/rGo nanostructured materials are also promising for various applications. Regarding the fabrication techniques, both plasma-based/electric field-assisted methods and thermal-type technologies are promising and are currently under development (Table 1). Microwave and radio-frequency plasmas are among the most promising techniques, although simple, cheaper DC-current plasma systems are also efficient for the fabrication of various CuO nanostructures and metamaterials.

Electric transport in copper oxide nanostructures

Despite the great diversity of CuO-based nanomaterials, electric field- and electric current-related phenomena play very important roles in most of their applications. As a result, a deep knowledge of the properties of single nanostructures or building blocks is of primary importance for understanding the fundamental issues of the device and its final performance. Unsurprisingly, many recently published reports discuss the electric field- and electric current-related phenomena in CuO-based nanomaterials. Recently, Kajli *et al.*¹⁸⁸ reported anomalies in the current density of copper oxide nanowires grown *via* a thermal process, *i.e.*, the current was observed to decrease as a function of the nanowire diameter. As shown *via* Raman and photoluminescence characterization, this behavior can be traced to the presence of (hole) traps in thicker Cu_2O NWs that hinder charge transport in p-type CuO. Quantum confinement has been reported by Ghosh *et al.*¹⁸⁹ for chemically synthesized freestanding thin (about 3–4 nm) CuO nanosheets at room temperature. Due to quantum confinement effects, the nanosheets displayed a rather large band gap of about 2.1 eV, larger than that for bulk CuO. These results point to the usefulness of these materials in supercapacitor devices. Shariffar *et al.* developed a UV-visible photodetector, based on copper oxide thin films, that displays enhanced optoelectronic properties if the films are synthesized at higher temperatures.¹⁹⁰ Such behavior could be explained by the influence of the grain size on the photocurrent, dark current, photosensitivity and responsivity. Zhou *et al.* also confirmed that the nanomaterial morphology has a major effect on the observed properties and types of application.¹⁹¹ They found that CuO nanostrip sensors possess a larger surface area and faster electron transfer compared with CuO nanowires and/or microspheres. They also displayed the superior electrochemical sensing of glucose and tetrabromobisphenol. As a result, one may expect that the morphology of CuO nanomaterials can influence the performance of devices that are based on these nanomaterials.¹⁹²

Other applications of CuO-based nanostructures

CuO-based nanostructures could be promising for many advanced applications that are not currently mature, *e.g.*, for space exploration. Light, thin, microwave-absorbing materials could be prepared using carbon/CuO microfiber composites, as

Table 1 CuO-based nanomaterials discussed in detail in this review. The spectrum of CuO-based materials and their applications reported recently (mainly for the past two years) are presented. Abbreviations: RFPE – radio-frequency plasma-enabled; MAHT – microwave-assisted hydrothermal technology; MTO – modified thermal oxidation; ES – electrospinning; TP – thermal processing

Type of nanostructure	Fabrication process	Feature of process/product	Potential application	Ref.
Nanowires	RFPE	Uniform surface growth	Sensing	68
Butterfly-like nanostructures	MAHT	CuO sheets were polarized by microwaves and arranged	Sensing	69
Nanowires	Thermal-induced	Electric-field assisted growth	Building blocks for nanodevices	72
Nanowires	MTO	Higher humidity produced higher yields of NWS	Nano-electro-mechanical systems	73
Porous hollow nanotubes	ES + TP	Nanotubes embrace copper oxide nanoparticles	Gas sensors	75
Nanowires	Plasma	Catalyst-free, single-crystalline	Sensing, catalysis	74
Nanowires	Facile solution-phase	Monoclinic polycrystalline structure	Sensors for volatile organic compounds	77
Nanowires	Anodization in sodium bicarbonate	Partial self-healing suggested	Various	78
Nanowires	Facile biosynthesis	<i>Sapindus mukorossi</i> fruit extract as a biosurfactant	Electrochemical sensing of dopamine	79
Nanowires	Thermal oxidation	Grown directly on V-shaped microgroves	Heat and fluid transport	84
Nanowires	Chemical synthesis	Cu ₂ S nanoparticles evolved into CuO nanowires	Catalysis of water splitting	85
Nano-needle arrays	Magnetron sputtering + wet etching + annealing	Grown on commercial ceramic tube	Tube-like gas sensors	113
Nanowires-3D copper hybrid	Electroless deposition	Hybrid with gold nanoparticles and Cu ₂ O nanowire-3D copper	Various	128
Graphene-based oxide nanofiber	Electrospinning	Decorated by Cu-nanoflowers	Biosensor for glucose	164
Complex nanomaterial	DC plasma thermal reaction method	Uniformly decorated by CuO nanocrystallines	Non-enzymatic glucose sensor	171
CuO/graphene	Reduction in ascorbic acid	Enriched with oxygen functional groups and defective sites	Gas sensing	154
Porous copper oxide	Hydrothermal reaction, calcination	Porous copper oxide heterostructured arrays	High-performance batteries	176
Nanowires	Reduction, oxidation	Synergetic combination of nanowires and graphene sheets	High-performance supercapacitors	178
Cu ₂ O nanoflakes	Facile electro-oxidation	Nanoporous Cu layer on flexible Cu foil	High-performance supercapacitors	185
Porous mesh	Thermal annealing	Activated/reduced graphene oxide on porous Cu	High-performance supercapacitors	186
Microfibers	Thermal annealing	C/CuO microfiber composites	Microwave absorbance	193
CuO/Cu ₂ O nano-needles	Electrophoretic deposition	Hierarchical porous distribution	High-performance supercapacitors	187

was demonstrated recently (Fig. 32).¹⁹³ Protective materials of this type would be indispensable for human protection in space and in other harsh environments. Moreover, CuO-based

nanostructures could be a base for cold electron emitters, capable of enhancing the characteristics of space propulsion systems.^{194–197}

Apart from the energy, sensing and catalysis applications discussed in this review, biomedical applications are among the most important fields that directly influence human health and wellbeing. Low-dimensional CuO-based nanostructures are quite promising for many biomedical applications that include, but are not limited to, *e.g.*, glucose sensing, H₂O₂ sensing, immunosensing, dopamine sensing, targeted cancer therapy, microbial warfare agents, wound healing and others (Fig. 33). While these applications are outside the scope of our review, we stress here in the Outlook the importance of biomedical applications, with a view to promoting further research activities in the field of bioactive low-dimensional copper oxide materials.

Future trends and tasks

Recently (2020), a quite detailed and comprehensive review on CuO-based nanomaterials for gas sensing was presented, where

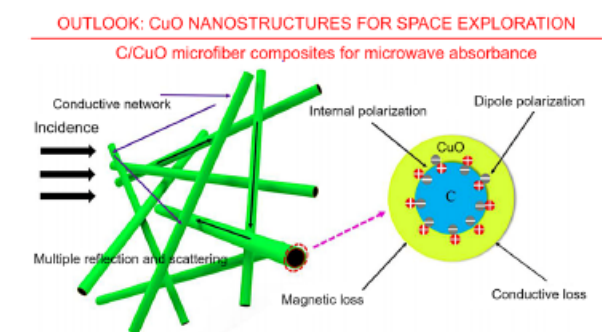


Fig. 32 Novel materials based on CuO nanostructures for possible space exploration applications. Mechanism of microwave energy absorbance on C/CuO microfiber composites. Reprinted with permission from Wei *et al.*, ref. 193. Copyright 2021, Springer.

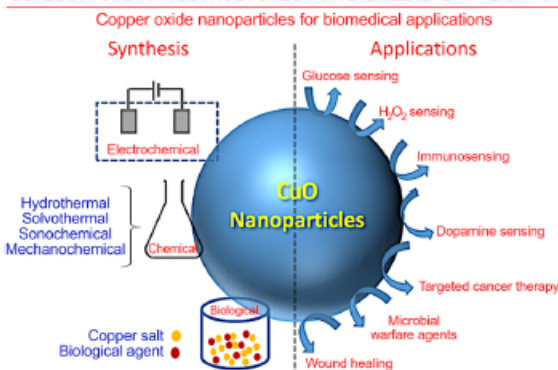


Fig. 33 Copper oxide nanoparticles for future biomedical applications. Low-dimensional CuO-based nanostructures are quite promising for many biomedical applications that include, but are not limited to, e.g., glucose sensing, H₂O₂ sensing, immunosensing, dopamine sensing, targeted cancer therapy, microbial warfare agents, wound healing and others. Reprinted with permission from Verma *et al.*, ref. 198. Copyright 2019, ACS.

several points were outlined as directions for the future research.⁶⁹ Based on newly achieved results and the broader scope of our review, which outlines energy applications and various techniques for the fabrication of CuO-based nanomaterials, we further develop and widen here the points suggested by Li *et al.*:

- ✓ loading CuO-based nanomaterials and nanostructures with noble metals and multiple doping with various elements is promising for many applications, and thus requires more research and design efforts;

- ✓ the interaction of CuO-based nanomaterials with various types of plasma is very important to deepen our understanding of the processes during the synthesis of various nano- and metamaterials;

- ✓ techniques for fabricating CuO-based nanomaterials in plasmas of various types should be further improved;

- ✓ in view of their relative simplicity, chemical and thermal methods for the fabrication of CuO-based nanomaterials are also very promising and should be further developed;

- ✓ further progress is required for understanding the routes and driving forces of the low-dimensional growth of nanomaterials;

- ✓ numerical simulations, in particular the artificial intelligence-based approaches, should be developed to ensure the deterministic design of CuO-based nanomaterials.

Conclusions

In this review paper we have outlined the most recent progress (mostly within the past 2 years) in the synthesis and application of low-dimensional copper oxide nanomaterials and complex materials designed on the basis of CuO. Further directions are discussed, and the most pressing problems outlined.

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