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# Optimizing the Cu<sup>2+</sup> ions and carbon-related defect centers ratio in g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu nanocomposites for supercapacitor applications<sup>†</sup>

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In this work, we synthesised a composite material based on g-C<sub>3</sub>N<sub>4</sub> as a carbon source, and ZnO doped with different Cu ion concentrations (ZnO:Cu x, x = 0, 0.2, 0.4, 0.6, 0.8, and 1%) using multiple synthesis approaches. The synthesized g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu nanocomposites were characterised using advanced techniques like X-ray diffraction, scanning and transmission electron microscopy, Fourier transform infrared spectroscopy, and UV-Vis adsorption and photoluminescence spectroscopy to evaluate the formation of the composite structure and the role of the dopant on the morpho-structural, optical, and compositional properties. Electron paramagnetic resonance and X-ray photoemission spectroscopy were employed to determine the carbon-related defect structure (V<sub>C</sub>) of the composite materials, which also revealed the presence of Cu ions in the ZnO lattice with a 2+ oxidation state. Electrochemical measurements, including galvanostatic charge-discharge and cycling stability tests, confirmed superior charge storage capacity, efficient electron transfer, and long-term durability. Notably, the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8-based supercapacitor device exhibited excellent cycling stability with 110% capacitance retention after 10,000 cycles, along with the highest power (917.81 W/kg) and energy density (25.24 Wh/kg), making it a promising candidate for high-performance energy storage applications. The results indicate that an increase in Cu concentration facilitates charge transport. The Cu-doped ZnO is primarily responsible for the pseudocapacitance mechanism in the supercapacitor device. Increasing the Cu doping concentration facilitates the appearance of redox reactions. However, once a certain dopant threshold is reached, any further increase in dopant concentration results in a sudden drop in current response, showing that the Cu<sup>2+</sup>/V<sub>C</sub> ratio plays an important role in the charge storage mechanism of the composites. Fine-tuning this ratio is a key parameter that has to be taken into account when designing carbon-based/metal oxides composite materials.

## 1 Introduction

As the global economy grows and fossil fuels are depleted, the research and commercialization of sustainable renewable energy sources and related technologies have become critical worldwide<sup>1,2</sup>. As a result, various energy-storage devices were invented<sup>3</sup>. Classical energy storage systems such as capacitors and

batteries continue to be used successfully. More recently, supercapacitors (SCs) are gaining popularity due to their greatly enhanced features, including high specific capacitance, high power density, fast charge-discharge time, and long life cycle<sup>4,5</sup>.

Supercapacitors are usually fabricated with an electrode material, electrolyte, and a separator. Depending on the electrode material, there are three main types of SCs: electric double-layer capacitors (EDLCs), pseudocapacitors, and hybrid supercapacitors<sup>1,6</sup>. Each device stores energy through distinctive mechanisms<sup>7-9</sup>, providing different outcomes dependent on the structure, composition, and stability of the applied electrode material<sup>8,10-14</sup>.

EDLC SCs store the energy electrostatically, not by chemical reactions, and rely on forming an electric double layer at the interface between the solid electrode and the liquid electrolyte when a voltage is applied<sup>7</sup>. The electrodes usually consist of high sur-

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face area carbon-based materials, such as activated carbon<sup>10</sup>, g-C<sub>3</sub>N<sub>4</sub><sup>11</sup>, carbon nanotubes<sup>10</sup>, graphene<sup>12</sup>, graphene oxide<sup>13</sup>, or reduced graphene oxide<sup>12</sup>. It has the disadvantage of a low energy density, between 5 and 15 Wh/kg, lower than other energy storage systems, such as Li-ion batteries (200 - 300 Wh/kg), Li-air batteries (11140 Wh/kg), flow batteries (10 - 50 Wh/kg) or fuel cells (800 - 10000 Wh/kg)<sup>14</sup>.

Pseudocapacitors have attracted significant scientific interest due to their high specific capacitance and their potential integration into portable, automotive, and industrial energy systems. The development of pseudocapacitor electrodes has advanced from the early application of metal oxides RuO<sub>2</sub>, MnO<sub>2</sub>, and ZnO to the advancement of conducting polymers such as polyaniline and polypyrrole<sup>15,16</sup>.

Among the metal oxides used in SCs, zinc oxide is a critical, low-cost, and eco-friendly II-VI group semiconductor material with good electrochemical properties<sup>17,18</sup>. Furthermore, during the ZnO synthesis, a highly defective structure such as zinc vacancies (V<sub>Zn</sub>), oxygen vacancies (V<sub>O</sub>), zinc interstitials (Zn<sub>i</sub>), and oxygen interstitials (O<sub>i</sub>) can be obtained, which can be investigated and guided in such a way that it may enhance its electrochemical performance<sup>19,20</sup>. Moreover, the concentration and type of defects within metal oxides can be effectively tailored by intentionally incorporating dopant species during doping, enabling precise control over the material's structural and electronic properties. Metals such as Cu<sup>21</sup>, Mo<sup>22</sup>, Fe<sup>23</sup>, Li<sup>24</sup>, or V<sup>22</sup> used in doping processes can further enhance their supercapacitive behaviour.

Hybrid SCs combine an electric double-layer capacitor with a pseudocapacitor in a single device, thus combining the high energy density of batteries with the high power density and long cycle life of supercapacitors<sup>9</sup>. Recently, carbon-based materials functionalized with metal oxides have been employed to enhance conductivity and surface area<sup>15</sup>. When metal oxides are decorated with carbon-based materials, several structural, interfacial, and electronic defects can form, like oxygen vacancies, interfacial dislocations, charge transfer, and carbon lattice defects<sup>25</sup>. Thus, defect engineering at the oxide-carbon interface is a key strategy for optimizing the performance of hybrid electrodes in supercapacitors<sup>23,24,26</sup>.

When combined with carbonaceous materials like GO, rGO, or g-C<sub>3</sub>N<sub>4</sub>, metal oxides provide an enhanced electron transfer path, high stability, excellent mechanical strength, and increased specific surface area<sup>27,28</sup>. The metal oxide contributes to pseudocapacitance through rapid and reversible redox reactions, while the carbon matrix stores charge via electric double-layer formation at the electrode-electrolyte interface. The intense interfacial contact between the two phases facilitates efficient charge transfer and ion diffusion, and the carbon's high surface area improves the utilization of redox-active sites. As a result, the hybrid electrode exhibits synergistic energy storage behavior, achieving both high energy density and excellent rate capability<sup>29,30</sup>.

Various metal oxides and carbon-based materials were examined as electrode materials for supercapacitor applications. Several studies with nanocomposites decorated with metal oxides such as SnO<sub>2</sub><sup>31</sup>, MnO<sub>2</sub><sup>32</sup>, Co<sub>3</sub>O<sub>4</sub><sup>33</sup>, RuO<sub>2</sub><sup>34</sup>, are hybridized with

rGO and the resultant performance composites are quite interesting, because they have tailor-made properties with enhanced performance compared to the individual materials<sup>35</sup>.

Wang *et al.*<sup>31</sup> studied the supercapacitive performance of the as-prepared SnO<sub>2</sub>, rGO, and SnO<sub>2</sub>-graphene nanocomposite electrodes using the CV technique. The CV curves of SnO<sub>2</sub>, rGO, and SnO<sub>2</sub>-graphene nanocomposite electrodes were recorded at a scan rate of 10 mV/s within a potential window of 0-1 V. The SnO<sub>2</sub>-graphene nanocomposite electrode showed a specific capacitance of 347.3 F/g, which is significantly higher than that of rGO (92.0 F/g) and SnO<sub>2</sub> (56.4 F/g) under identical conditions, highlighting its superior capacitance and more efficient charge propagation within the electrode.

MnO<sub>2</sub> decorated on carbon structures and used as an electrode material for supercapacitors exhibited remarkable improvements in electrochemical properties, as reported by Ramezani *et al.*<sup>32</sup>. The pure MnO<sub>2</sub> achieved a specific capacitance of 55.7 F/g, while MnO<sub>2</sub>-carbon nanotubes with 180 F/g, and MnO<sub>2</sub>-graphene nanosheets with 310 F/g markedly surpassed it. Moreover, a ternary composite using two carbon structure, MnO<sub>2</sub>-graphene nanosheets-carbon nanotubes composite achieved a specific capacitance of 367 F/g at the same scan rate of 20 mV/s, greater than all the previous values, emphasizing the synergistic integration of graphene nanosheets and carbon nanotubes, which collectively promote efficient electrolyte ion transport and reinforce the mechanical robustness of the electrode structure<sup>32</sup>.

In a study that compared the performances of ZnO and g-C<sub>3</sub>N<sub>4</sub>-ZnO nanocomposite using urea and thiourea precursors, as electrode material, it was found that g-C<sub>3</sub>N<sub>4</sub>-ZnO using an urea precursor and g-C<sub>3</sub>N<sub>4</sub>-ZnO using a thiourea precursor exhibited higher specific capacitance of 415 and 408 F/g compared to pure ZnO 223 F/g with better cycling stability, indicating improved redox activity and superior charge storage ability<sup>36</sup>.

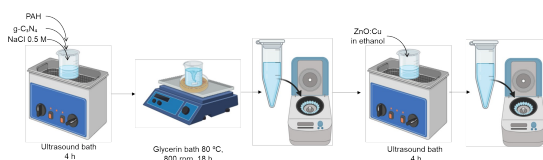
The present study focuses on advancements in supercapacitors through the development of hybrid electrode materials. Here, an innovative composite comprising g-C<sub>3</sub>N<sub>4</sub> was designed as a carbon source, and copper-doped ZnO nanoflowers were synthesized and evaluated to demonstrate their supercapacitor performance. The nanocomposite presents two distinct paramagnetic centers: the Cu<sup>2+</sup> ions and carbon-related defect centers (V<sub>C</sub>). The spectral features associated with Cu<sup>2+</sup> exhibit noticeable changes with increasing doping concentration, indicating a direct correlation between copper incorporation and the resulting magnetic environment, as evidenced by EPR spectroscopy. In contrast, the signal attributed to the C-related defect centers remains essentially constant, suggesting that these intrinsic defects are not significantly influenced by the level of Cu doping<sup>37,38</sup>. The relative contribution of each paramagnetic species, expressed as the ratio of their EPR signals and normalized by linewidth and intensity, constitutes a key analytical parameter. This ratio provides valuable insight into the structural and electronic modifications induced by doping and is closely linked to the electrochemical behavior of the resulting material. Consequently, it serves as an important indicator for assessing and optimizing the nanocomposite's performance when employed as an active material in supercapacitor devices. Our investigations align with supercapac-



itor challenges regarding materials fabrication, cost, scalability, and limited energy storage.

## 2 Materials and methods

### 2.1 Chemicals



Scheme 1 Schematic representation of the synthesis procedure used to synthesize g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu

The reagents and solvents used in this research were of analytical grade and were used without any supplementary purification. The following reagents and solvents were used in the synthesis procedure: urea - CH<sub>4</sub>N<sub>2</sub>O (ACS Reag. Ph., VWR Chemicals) was used as a carbon source and precursor for obtaining of g-C<sub>3</sub>N<sub>4</sub>, sodium hydroxide - NaOH (Sigma Aldrich) was the precipitating agent in the ZnO synthesis, zinc nitrate hexahydrate Zn(NO<sub>3</sub>)<sub>2</sub> · 6H<sub>2</sub>O (Alpha Aesar) was the zinc source, copper nitrate - Cu(NO<sub>3</sub>)<sub>2</sub> (Alpha Aesar) was used to dope the ZnO with Cu ions, sodium chloride - NaCl (ACS Reag. Ph., VWR Chemicals) controls the aggregation and distribution of the nanoparticles, poly-allylamine hydrochloride - PAH (Alfa Aesar) was the linking polymer between ZnO:Cu and the carbon source. The aqueous solutions were prepared from Milli-Q water produced by a Direct-Q 3UV system (Millipore, Bedford, MA).

### 2.2 Samples preparation

ZnO:Cu nanoparticles, having a nano-flower shape, were prepared through a chemical precipitation method as depicted in Scheme 1 where 10 mM of zinc nitrate hexahydrate was dissolved in 100 ml of distilled water under magnetic stirring, and NaOH 3 M was added until a pH of 12 was reached, followed by stirring on an orbital shaker for 4 h. The procedure is described in detail in a previous article by Stefan *et al.*<sup>39</sup>.

Doping ZnO with Cu is typically kept below 1% because the solubility of Cu<sup>2+</sup> in the ZnO lattice is very limited. Higher concentrations lead to the segregation of secondary phases such as CuO, Cu<sub>2</sub>O, or metallic Cu, which distort the wurtzite structure, introduce significant lattice strain, and create defect states that act as non-radiative recombination centers. These effects degrade the optical, electrical, and catalytic performance of the material<sup>40</sup>. At low concentrations (<1%), Cu can substitute Zn sites without severely perturbing the crystal lattice, enabling controlled modification of electronic states and enhanced visible-light absorption. However, excessive Cu doping increases Jahn–Teller distortions and defect formation, reduces carrier mobility, and shifts the material's behavior from intrinsic ZnO properties to those dominated by Cu-based phases<sup>41,42</sup>. Thus, sub-percent Cu doping offers the best balance between structural stability and functional improvement.

g-C<sub>3</sub>N<sub>4</sub> was obtained by the thermal decomposition of urea,

heating it at 550 °C for 2 h as depicted in Scheme 2. The g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu nanocomposites were assembled from the two individual components using the linking polymer polyallylamine hydrochloride (PAH). The volumetric ratio between g-C<sub>3</sub>N<sub>4</sub> and ZnO:Cu was 1:1.

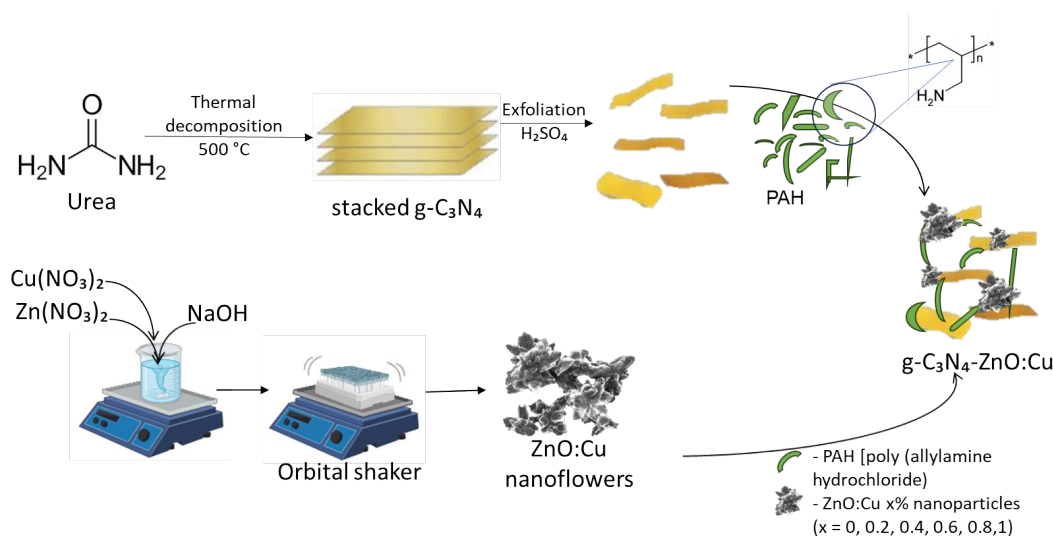
A stock solution of 500 mL saline solution NaCl 0.5M was prepared by weighing 14.6 g solid NaCl and adding water until 500 mL, with 2-3 minutes stirring until completely dissolved. g-C<sub>3</sub>N<sub>4</sub> was uniformly dispersed in the saline solution of PAH obtained by dissolving 0,7 g PAH in 100 mL NaCl 0.5 M by sonication for 4 h. The reaction mixture was heated to 80 °C for 18 h with continuous stirring, then cooled to room temperature. After centrifugation, the remaining material was added to an ethanol suspension of the ZnO:Cu nanoparticles and ultrasonicated for an additional 4 h. The mixture was centrifuged again, and the obtained precipitate was washed and dried at 65 °C. The entire synthesis procedure is summarized in Scheme 2.

### 2.3 Morpho-structural characterization methods

X-ray diffraction (XRD) measurements were performed on a Smart Lab Rigaku diffractometer with Cu-K $\alpha$  radiation ( $\lambda = 1.54$  Å). All measurements were conducted at room temperature over a  $2\theta$  range of 10-90°, with a 0.01° step. The nanocomposites' size, shape, clustering, and particle size distributions were examined with Scanning Transmission Electron Microscopy (STEM) on a Hitachi HD2700 electron microscope. Photoluminescence spectra (PL) were acquired on an FS5 spectrofluorometer (Edinburgh Instruments) equipped with a 150 W CW ozone-free Xenon arc lamp, a Czerny-Turner with plane grating monochromator, and a PMT-900 emission detector. FTIR spectra were obtained in the 400–4000 cm<sup>−1</sup> spectral range with a JASCO 6100 FTIR spectrometer using the KBr pellet technique. The spectral resolution to record the IR spectra was 2 cm<sup>−1</sup>. JASCO V570 UV-VIS-NIR spectrophotometer equipped with absolute reflectivity measurement, JASCO ARN-475 accessory was used to measure the absorption spectra. The UV-Vis absorption properties of the synthesized samples were investigated over the 200-800 nm spectral range. Raman measurements on all composites were performed using a Renishaw InVia Raman system paired with a Leica microscope. The excitation source was a near-infrared (NIR) diode laser at 830 nm. The Raman measurements covered a wavenumber range of 700-1650 cm<sup>−1</sup> with a spectral resolution of 1 cm<sup>−1</sup>. Experiments used a 1 s laser exposure time and a single acquisition at 30 mW laser power. XPS qualitative and quantitative composition of the samples was examined with a custom-built SPECS spectrometer using a Mg anode (1253.64 eV) as the X-ray source. The samples were cast onto a sample holder. Spectra were analyzed with the CASA software using relative sensitivities, transmission factors, and electronic mean-free-path factors. Electron paramagnetic resonance (EPR) spectroscopy was performed at room temperature using a 9.88 GHz X-band Bruker E-500 ELEXSYS spectrometer. Nitrogen adsorption-desorption isotherms were recorded at 77 K using a BelSorp MaxX equipment (Microtrac BEL Corporation, Japan). Samples pretreatment consisted of degassing under vacuum at 80°C for 6 h before N<sub>2</sub>







Scheme 2 Illustrative representation of the procedure used to synthesize g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu

sorption. The standard BET method was used to calculate the specific surface area ( $p/p_0$  in the range of 0.05–0.25), total pore volume was estimated at  $p/p_0 = 0.95$ , and the pore size distribution was analyzed with the Barrett-Joyner-Halenda (BJH) method from the N<sub>2</sub> desorption branch.

## 2.4 Electrical properties measurements

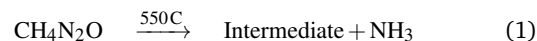
Symmetrical all-in-one supercapacitor devices were assembled using the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu composites as an electrode material (3 mg) while a 6 M KOH aqueous solution served as an electrolyte (100  $\mu$ L). A glass fiber separator was employed between the electrodes. A circular-shaped stainless-steel bolt on both sides of the device serves as a current collector. The areal mass loading of the electrodes on this device is approximately 2.65 mg/cm<sup>2</sup>, with a total mass of 3 mg per electrode. As the powder sample is directly loaded onto the current collector without a thin film or substrate, the focus is on measuring the performance of the synthesized sample; the electrode thickness is not measured. The focus remains on maintaining consistent mass loading across all devices. In the symmetric device, electrochemical testing was performed without any additional booster material, such as carbon, and only the mass of the single electrode was used to calculate specific capacitance and Ragone plot values to report the contribution from the individual electrode. A BioLogic VMP 300 electrochemical workstation was used to test the electrochemical performance of the assembled supercapacitor devices. The techniques included cyclic voltammetry (CV), potentiostatic electrochemical impedance spectroscopy (PEIS), and galvanostatic cycling with potential limitation (GCPL). CV measurements were performed within a voltage window of 0 to 1 V, with scan rates ranging from 2 to 200 mV/s. PEIS was performed with an AC signal amplitude of 10 mV across a frequency range of 10 mHz to 1 MHz. GCPL tests were conducted at a current density of 0.8 A/g, with the potential cycled over a 0–1 V window at a scan rate of 10 mV/s. Specific capacitance, energy density, and power density

were calculated based on the data from CV and GCPL as described in previous studies<sup>24,39</sup>.

## 3 Results and discussion

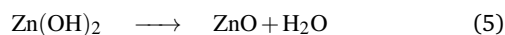
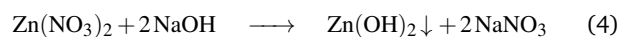
The general criteria for high electrochemical performance are chemical stability and surface features<sup>43</sup>. These material-specific characteristics can be developed using appropriate synthesis procedures and by carefully optimizing the experimental parameters. Considering the variables above, g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cux nanocomposites were prepared by combining previously synthesized individual components at the same volume ratio. The heterojunction was realized using poly(allylamine hydrochloride) (PAH) as a linking polymer. PAH is a cationic polymer with primary amine groups, making it highly water-soluble and reactive<sup>44</sup>. Because of its strong electrostatic interactions, it is often used to synthesize composite materials.

Thus, g-C<sub>3</sub>N<sub>4</sub> was formed by thermal condensation followed by polymerisation of urea at around 550 °C. The obtained bulk g-C<sub>3</sub>N<sub>4</sub> was ground into fine powder, followed by chemical exfoliation with H<sub>2</sub>SO<sub>4</sub> to obtain g-C<sub>3</sub>N<sub>4</sub> nanosheets. The general chemical reactions for thermal decomposition of urea are described in the following order<sup>45,46</sup>:



In a separate synthesis procedure, copper-doped ZnO nanoflowers were obtained by chemical precipitating of zinc nitrate in sodium hydroxide as described in Equations 4 and 5.





The g-C<sub>3</sub>N<sub>4</sub> nanosheets were functionalised with PAH to increase their reactivity. The newly existing functional groups contribute to forming new bonds in the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu composites. Scheme 2 outlines the synthesis procedures for obtaining g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu<sub>x</sub> nanocomposites.

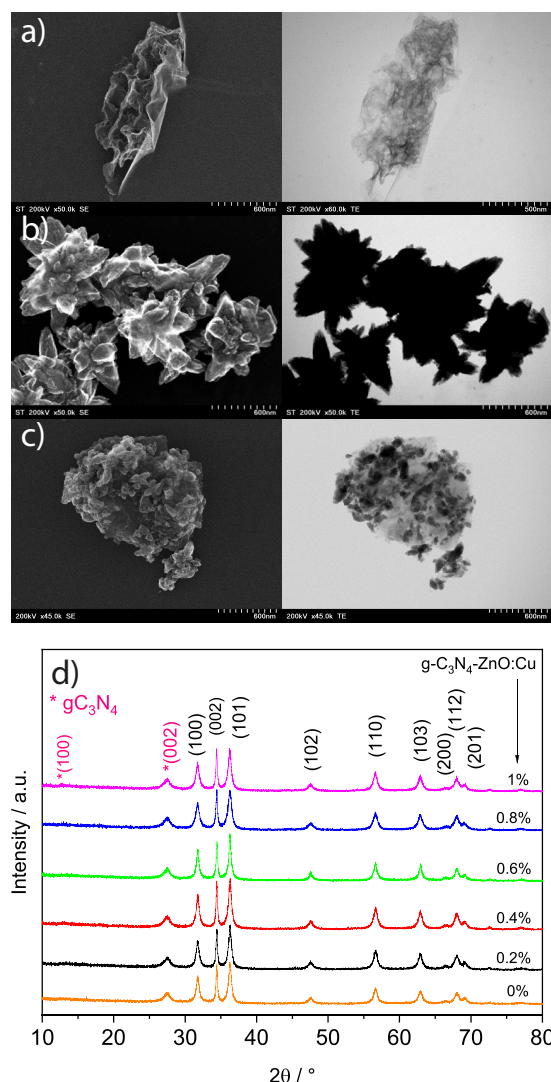


Fig. 1 SEM/TEM images of (a) g-C<sub>3</sub>N<sub>4</sub>, (b) ZnO nanoflowers, (c) g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.4 and (d) X-ray diffraction patterns of the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu composites.

The morphology of g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu was investigated by SEM and TEM microscopy, as shown in Figure 1a-c. As an example, in Figures 1a and 1b the images corresponding to pristine g-C<sub>3</sub>N<sub>4</sub> and, respectively, ZnO nanoflowers are depicted, as well as a representative image of the composites, namely g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.4 (Figure 1c). The TEM and SEM images in Figure 1a show the presence of thin sheets, a morphology specific to g-C<sub>3</sub>N<sub>4</sub>. From the TEM and SEM images presented in Figure 1b, the morphol-

ogy of ZnO nanoparticles is revealed as elongated petal-like particles. The mean size of the ZnO:Cu particles is approximately 120 nm. In the case of the composite materials (Figure 1c), it can be observed that several ZnO nanoparticles are firmly attached to the g-C<sub>3</sub>N<sub>4</sub> thin sheets. It should be noted that comparable morphologies were obtained for all processed samples.

The elemental analysis, as indicated by EDS mapping measurements results presented in Figure S-EDS<sup>†</sup>, reveals the presence of Zn, O, Cu, C, and N, confirming the assembly of ZnO:Cu nanoparticles attached to thin g-C<sub>3</sub>N<sub>4</sub> sheets, and the presence of the Cu dopant ions distributed uniformly in the host ZnO lattice.

XRD diffraction was performed to identify the structural crystalline phases and the sample's structure. Figure 1d shows the diffraction patterns of g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu composites. As observed, the diffractograms contain diffraction patterns specific to both composite material components. The peaks observed at  $2\theta = 12.71^\circ$  and  $27.47^\circ$  are indexed to (100) and (002) corresponding to the planes of g-C<sub>3</sub>N<sub>4</sub> (card JCPDS 79-0208), while all the other peaks correspond to the hexagonal wurtzite structure of ZnO (space group P6<sub>3</sub>mc, JCPDS 79-0208). No other crystalline phases are present, showing the samples' purity. The cell parameters were calculated using Rietveld refinement and are summarized in Table 1. Cu doping causes a small increase in cell volume up to 0.8 mol%, followed by a substantial drop at the higher Cu concentrations (1 mol%). Even though Cu<sup>2+</sup> ions have an ionic radius (0.73 Å) similar to that of Zn<sup>2+</sup> ions (0.74 Å), an increase in the unit cell volume is observed. This behavior can be explained by the fact that some Cu<sup>2+</sup> ions partially replace Zn<sup>2+</sup> ions, while others occupy interstitial sites. Interstitial positions can give rise to various lattice defects, such as Zn interstitials or oxygen vacancies. At a Cu doping concentration of 1 mol%, a significant reduction in the cell volume is observed, probably due to the segregation of Cu<sup>2+</sup> ions on the surface of ZnO and the formation of zinc vacancies. The average crystallite size, calculated by the Williamson-Hall method, ranges from 11 to 14 nm.

FT-IR spectroscopy was used to investigate the functional groups and chemical bonds in the samples. Figure 2a shows the FT-IR spectra of g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu1 composite in the range of 350 - 2000 cm<sup>-1</sup>. At 412 cm<sup>-1</sup>, a broad and intense band corresponding to the stretching mode of the Zn-O bond<sup>47</sup>. The peak centered at 807 cm<sup>-1</sup> is associated with the triazine units, while the peaks at 1235 cm<sup>-1</sup> and 1637 cm<sup>-1</sup> correspond to the C-N and C=N stretching vibration modes of g-C<sub>3</sub>N<sub>4</sub><sup>48</sup>. The peak at 1315 cm<sup>-1</sup> is assigned to C-C and C-N bending vibrations, whereas the band at 1402 cm<sup>-1</sup> corresponds to the heptazine-derived repeating units<sup>49</sup>. A shift to lower wavenumbers of the g-C<sub>3</sub>N<sub>4</sub>-specific peaks by Cu doping of ZnO is observed (see Figure S-FT-IR<sup>†</sup>). This shift indicates that the conjugated system of g-C<sub>3</sub>N<sub>4</sub> was stretched, and a broadly conjugated system containing g-C<sub>3</sub>N<sub>4</sub> and ZnO had formed<sup>50</sup>.

UV-Vis spectroscopy was employed to investigate the optical properties of g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu samples, and the results are shown in Figure S-UV-Vis<sup>†</sup>. All samples exhibit significant absorption in the 200 - 400 nm range, consistent with the characteristic behavior of ZnO<sup>51</sup>, with a maximum absorption at 366 nm corresponding to interband transitions<sup>39</sup>. Increasing the copper concentra-



Table 1 Cell parameters, cell volume, and crystallite mean size of the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu<sub>x</sub> composites, obtained from the Rietveld analysis of the XRD data, and BET surface area and total pore volume of the composites obtained from the nitrogen adsorption-desorption isotherms.

Sample	ZnO:Cu				g-C <sub>3</sub> N <sub>4</sub>			$S_{\text{BET}}$ / m <sup>2</sup> /g	$V_p$ / cm <sup>3</sup> /g
	$a = b$ / Å	$c$ / Å	$V$ / Å <sup>3</sup>	$D$ / nm	$a = b$ / Å	$c$ / Å	$V$ / Å <sup>3</sup>		
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0	3.243	5.199	47.37	12	6.492	2.516	91.8	16.15	0.043
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.2	3.243	5.202	47.39	11	6.492	2.516	91.8	16.19	0.042
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.4	3.246	5.196	47.39	12	6.492	2.516	91.8	19.84	0.056
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.6	3.247	5.196	47.40	14	6.492	2.516	91.8	19.02	0.051
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.8	3.243	5.199	47.36	10	6.492	2.516	91.8	21.80	0.063
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu1	3.241	5.195	47.26	13	6.492	2.516	91.8	20.61	0.057

tion shifts the absorption spectra towards higher wavelengths. The semiconductor's band gap energy ( $E_g$ ) was calculated based on the absorption spectrum using Tauc's equation with<sup>52</sup>:

$$\sqrt[n]{\alpha h\nu} = A(h\nu - E_g) \quad (6)$$

In this equation,  $\alpha$  is the absorption coefficient,  $h\nu$  is the photon energy,  $A$  is a constant, and  $n$  is the Tauc exponent, which depends on the type of electronic transition (e.g.,  $n = 2$  for indirect allowed transitions and  $n = 1/2$  for direct allowed transitions). To determine  $E_g$ ,  $\sqrt[n]{\alpha h\nu}$  was plotted as a function of  $h\nu$ , and the linear portion of the graph is extrapolated to the  $x$ -axis to find the intercept corresponding to the band gap energy.

Figure 2b shows the corresponding Tauc's plot with the band gap energy values for all samples, and a slight decrease is observed with increasing copper doping. The reduction in the band gap energy is attributed to several factors: the presence of Cu d-states below the conduction band<sup>53</sup>, the existence of a tensile stress induced by a mismatch between the dopant ions and Zn radius<sup>54</sup>, the formation of defect levels by doping within the band gap<sup>55</sup>, to the strong sp-d exchange interaction between the band electrons of ZnO and the localized d electrons of Cu<sup>2+</sup> ions substituting for Zn<sup>2+</sup> ions, which shifts the Fermi level into the valence band<sup>56</sup>. In the undoped state, the Fermi level is located at the center of the band gap, and its position depends on the type of semiconductor<sup>39</sup>. Doping with Cu ions modifies the semiconductor's electrical and optical properties by introducing acceptor levels near the valence band. These levels facilitate electron excitation from the valence band, increasing hole concentration and enhancing p-type conductivity. Copper has one fewer valence electron than zinc; that is why, when copper is introduced into zinc oxide, it acts as a p-type dopant and generates holes in the valence band<sup>57</sup>. As the hole concentration increases, the Fermi level shifts downward, approaching the valence band. There are more unoccupied states (holes) at lower energies, and as a result, the Fermi level decreases<sup>58</sup>. Cu doping often results in slightly reduced band gap energy due to the formation of localized states and increased electron-hole interactions. Similar behavior was reported by Ma *et al.*<sup>58</sup>, who observed that increasing the Cu dopant concentration leads to a decrease in the optical band gap of Cu-doped ZnO.

The textural properties of the samples were evaluated using the BET method and are summarized in Table 1 and Figure S-BET<sup>†</sup>. As expected, we observe that both the specific surface area and the pore volume are almost unchanged with the Cu doping con-

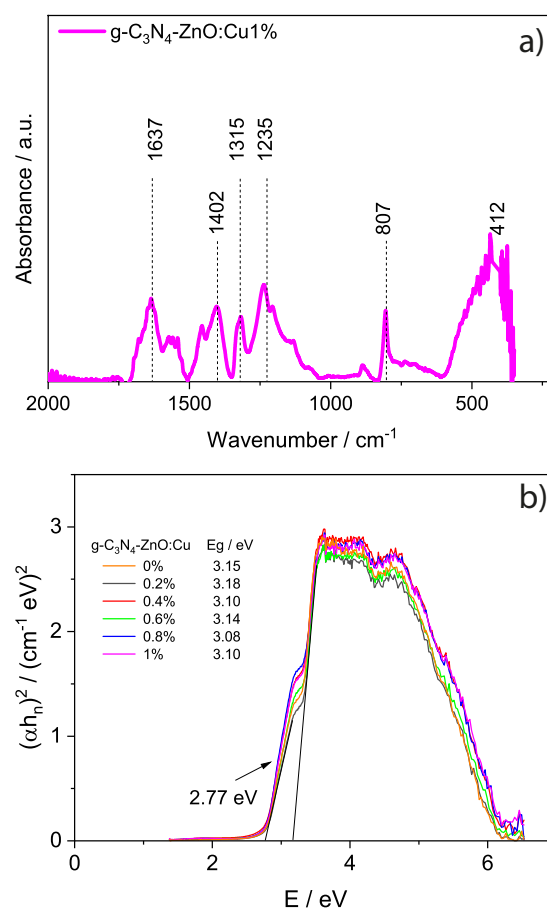


Fig. 2 (a) FT-IR spectroscopy for the sample g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu1 and (b) estimation of the band gap energy  $E_g$  based on Tauc's equation for all the synthesized samples.

centration. At a concentration of 0.8% Cu a maximum of 21.8 m<sup>2</sup>/g is obtained, after which a slight decrease is observed. In general, small variations in specific surface area can arise from the synthesis procedure. The specific surface area is a key factor for materials used as electrode materials for supercapacitor applications, but as evidenced by the BET measurements, Cu doping of ZnO in the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu composites is not influencing the specific surface area, indicating that the improvements in specific capacitance, which are described in the following, are due to other factors.

The Raman spectra of the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu samples have the characteristic vibrational signals of graphitic carbon nitride and



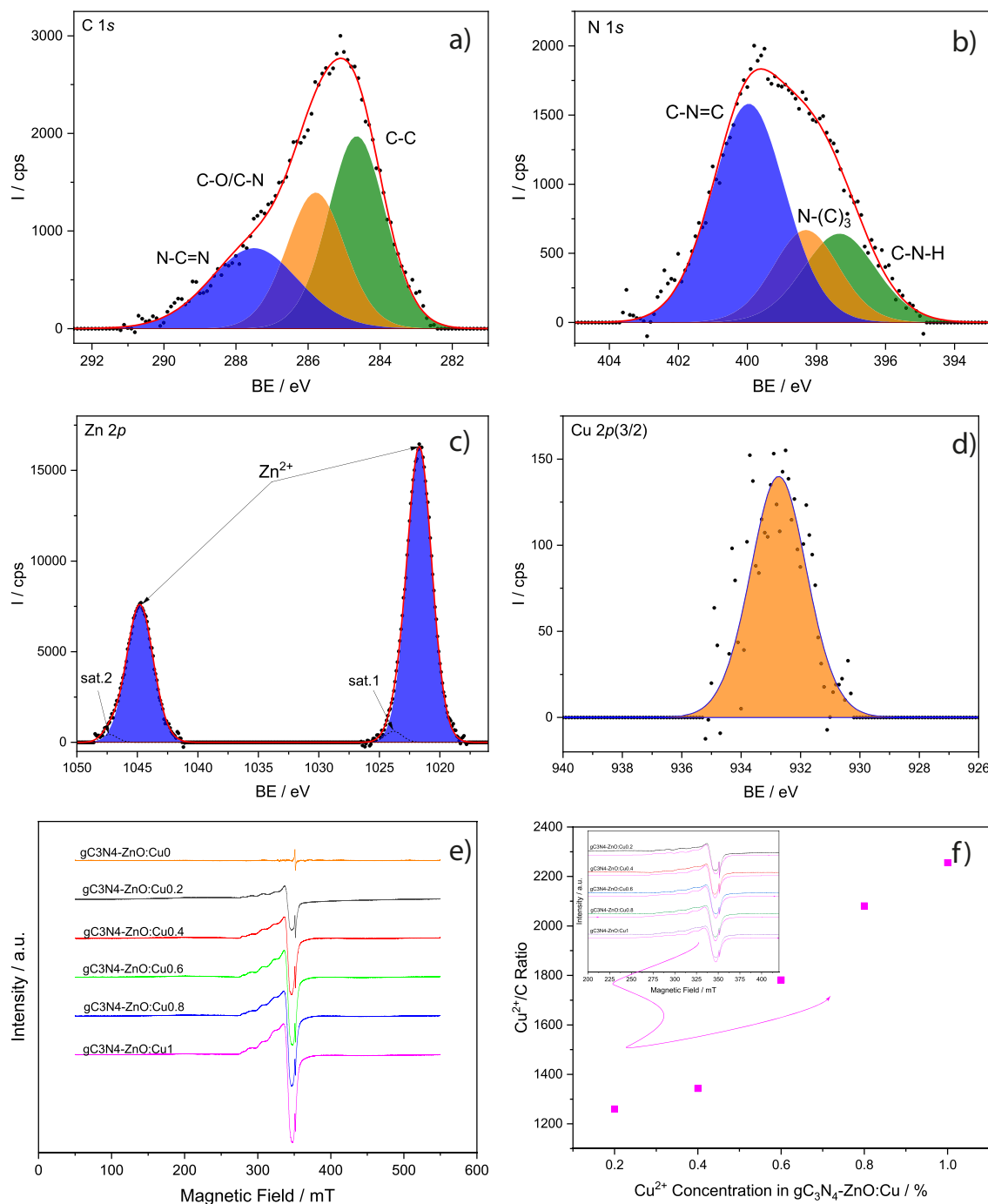


Fig. 3 XPS analysis of g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu<sub>0.8</sub>, as follows: a) C 1s core level, b) N 1s core level, c) Zn 2p core-level doublet, d) Cu 2p(3/2) core-level, and e) EPR spectra for all the composites while in f) the EPR signal ratio between the Cu<sup>2+</sup> ions and the carbon based defect centers, obtained from the simulations of the EPR spectra of the composites, is presented as a function of the doping Cu concentration. The inset of (f) shows the simulation results.

show minor differences between all Raman spectra. As shown in Figure S-Raman<sup>†</sup>, several peaks are intense. The band at 706 cm<sup>-1</sup> corresponds to the heptazine ring-breathing mode. The peaks at 751 and 767 cm<sup>-1</sup> are attributed to out-of-plane deformations of the heptazine units, while the band at 977 cm<sup>-1</sup> is assigned to an additional out-of-plane deformation of the heterocyclic rings. The vibrational modes at 1116 and 1152 cm<sup>-1</sup> originate from C–N stretching and C–N–c in-plane bending. The

1232 cm<sup>-1</sup> band is associated with C–N stretching. In the higher-frequency region, the bands at 1310 and 1350 cm<sup>-1</sup> correspond to defect-related (D band) modes and C–N/C=N stretching, and the peaks at 1405 and 1481 cm<sup>-1</sup> arise from C–N and C=N stretching vibrations. The bands at 1558 and 1620 cm<sup>-1</sup> fall within the graphite-like G-band region.<sup>59–61</sup> To evaluate defect concentration, the ratio of the intensity of the D region (1350 cm<sup>-1</sup>) to the G region (1558 cm<sup>-1</sup>) was calculated for each sam-





Sample	Cu <sup>2+</sup>					V <sub>C</sub> g <sub>C</sub>	Weight Cu <sup>2+</sup> /V <sub>C</sub>
	g <sub>  </sub>	g <sub>⊥</sub>	A <sub>  </sub> / MHz	g <sub>strain</sub>	A <sub>strain</sub>		
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.2	2.355	2.0817	449	0	0	2.0109	1260
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.4	2.355	2.0817	449	0.07	80	2.0109	1343
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.6	2.355	2.0817	449	0.08	110	2.0109	1782
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.8	2.355	2.0817	449	0.095	140	2.0109	2081
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu1	2.355	2.0817	449	0.11	169	2.0109	2256

Table 2 Main spin system parameters used to simulate the EPR spectra of the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu composites: one spin system for Cu<sup>2+</sup> centers and the other one for C-related defects (V<sub>C</sub>), as well as the ratio between them

ple. A higher  $I_D/I_G$  ratio indicates a greater degree of structural disorder<sup>62,63</sup>. The obtained values are as follows: undoped g-C<sub>3</sub>N<sub>4</sub>-ZnO - 0.691, 0.2% Cu - 0.642, 0.4% Cu - 0.690, 0.6% Cu - 0.697, 0.8% Cu - 0.710, and 1% Cu - 0.663. These results show that the 0.8% Cu-doped sample exhibits the highest  $I_D/I_G$  ratio, suggesting that this composition has the greatest defect concentration.

Further information on the material's structure, defects, and impurities was obtained through photoluminescence spectroscopy. The PL emission spectra were recorded at an excitation wavelength of 325 nm and are similar for all g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu samples, as depicted in Figure S-PL<sup>†</sup>. A broad emission band spanning 400-750 nm is observed. Because the broad response spanned a wide wavelength range, the spectra were deconvoluted with a Gaussian fit. This operation helps us identify the transitions responsible for the emissions and determine the positions of each spectral peak. The results are shown in Figure S-PLdeconv<sup>†</sup> for g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu1 as an example. Also, the Gaussian peak fitting parameters ( $R^2$ , residuals) are provided in Table S-PL<sup>†</sup>. According to previous PL studies, the bandgap states of g-C<sub>3</sub>N<sub>4</sub> consist of an sp<sup>3</sup> C-N  $\sigma$  band, an sp<sup>2</sup> C-N  $\pi$  band, and the lone pair (LP) state of the bridging nitrogen atom. Thus, the emission band centered at 429 nm was assigned to the  $\pi^* \rightarrow \pi$  transition, the band at 451 nm to the  $\sigma^* \rightarrow$  LP transition, and the emission at 484 nm to the  $\pi^* \rightarrow$  LP transition<sup>64</sup>. Compared to the emissions of g-C<sub>3</sub>N<sub>4</sub>, the composite samples show an enhancement of these emissions due to the charge recombination process at the interface of the two components.

XPS qualitative analysis was performed by examining the survey spectrum obtained from the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8 sample. In Figure S-XPS<sup>†</sup>, the expected Zn, O, C, and N elements are clearly observed. In addition, traces of Cl are present, originating from the synthesis reagents. The Cu 2p lines are not observed in the survey due to the low concentration; however, in the narrow-scan region, as shown in Figure 3d, the data is quantifiable. Also, the expected Cu LMM Auger peak at BE = 334 eV is not marked because the more intense Zn LMM peak overlaps it.

XPS qualitative and quantitative composition was done for the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8 sample; the results are shown in Figure 3a-d. The Zn 2p core-level doublet XPS spectra are shown in Figure 3a. The doublet positioned at 1021.7 eV (2p<sub>3/2</sub>) and 1044.8 eV (2p<sub>1/2</sub>) is indicative of typical of Zn<sup>2+</sup> binding energy. Also, two satellite peaks at higher binding energies were used in the deconvolution. The Cu 2p(3/2) core-level XPS spectrum is shown in Figure 3b. The peak is at 932.7 eV, while the calculated Cu/Zn

ratio is 0.008, showing a good doping yield and the expected presence of Cu<sup>2+</sup> ions, with no additional oxidation states. The C 1s core level line spectrum is presented in Figure 3c. The deconvolution was performed by considering the specific lines of graphitic carbon nitride<sup>65</sup>. The binding energy positions are 284.7 eV for C-C, 285.8 eV for C-O/C-N, and 287.5 eV for N-C=N. In this case, the C-C peak has a higher relative intensity, as expected for g-C<sub>3</sub>N<sub>4</sub>, due to the overlapping contribution from ZnO and g-C<sub>3</sub>N<sub>4</sub> interaction<sup>66</sup>. The N 1s core level line spectrum is presented in Figure 3d. The deconvolution shows three peaks at 397.3, 398.3, and 400.0 eV, assigned to C-N=C, N-(C)<sub>3</sub>, and C-N-H groups, respectively. The XPS analysis confirmed the successful Cu doping of ZnO and an interaction between ZnO with g-C<sub>3</sub>N<sub>4</sub>.

The presence of Cu ions in the ZnO lattice and carbon-related defect centers in g-C<sub>3</sub>N<sub>4</sub> were confirmed by EPR spectroscopy. The samples containing Cu show an axial EPR signal specific to Cu<sup>2+</sup> ions in axial symmetry, with four parallel components centered at  $g_{||} \approx 2.36$  having a hyperfine constant  $A_{||} \approx 150$  G and one component at  $g_{\perp} \approx 2.07$  (see Figure 3e). The hyperfine structure is due to the interaction between the electronic spin  $S = 1/2$  of the Cu<sup>2+</sup> ion and the nuclear spin  $I = 3/2$  of the copper nuclei <sup>63</sup>Cu and <sup>65</sup>Cu<sup>67,68</sup>. Additionally, at  $g = 2.01$ , the EPR spectra show a sharp line due to the unpaired electron localized in the  $\pi$ -conjugated structure of g-C<sub>3</sub>N<sub>4</sub><sup>69</sup>.

The fact that  $g_{||} > g_{\perp} > g_e$  (where  $g_e = 2.0023$  is the  $g$  value for a free electron) suggests that the Cu<sup>2+</sup> ions are subject to an elongated tetragonal distortion<sup>70</sup>. Based on the Hamiltonian spin parameters, the covalent parameters for the in-plane  $\sigma$  bond can be evaluated using the following expression<sup>71</sup>:

$$\alpha^2 = \frac{A_{||}}{0.036} + (g_{||} - g_e) + \frac{3}{7}(g_{\perp} - g_e) + 0.04 \quad (7)$$

The obtained value for  $\alpha^2$  is 0.84. It is known that for  $\alpha^2 = 1$  the bond is fully ionic, and for  $\alpha^2 = 0.5$  it is fully covalent. Considering the above results, it is not possible to clearly distinguish between covalent and ionic character. However, larger  $\alpha^2$  values indicate that the in-plane  $\sigma$  bond is moderately ionic. As the concentration of Cu<sup>2+</sup> ions increases, the EPR signal intensifies. The entire EPR spectra can be visualized in Figure 3e.

To get further insights on the importance of the paramagnetic ions, Cu<sup>2+</sup> and C-related defect centers (V<sub>C</sub>), the EPR spectra were simulated using the Easyspin package<sup>72</sup>. As depicted in Table 2, two spin systems were used for the simulations. The first system describes the Cu<sup>2+</sup> centers, for which a spin 1/2 was em-



played with a hyperfine coupling. The broadening due to dipole-dipole interactions between the  $\text{Cu}^{2+}$  centers, or structural disorder due to the Cu-doping, were added to the system as so-called g- and A-strains, which are the line-broadening components of the g and A tensors. Dipole-dipole interactions are magnetic interactions between unpaired electron spins. In dilute solutions, this primarily leads to broadening of EPR spectral lines, with the line width proportional to the spin concentration, and even to splitting signals depending on their magnitude and orientation relative to the magnetic field. This interaction depends on the distance between the two spins and on the orientation of the vector connecting them with respect to the field vector<sup>68</sup>. Structural disorder induced by Cu doping can broaden the EPR lines due to variations in the local atomic environment around the paramagnetic centers. Doping can introduce local distortions, alter crystal symmetry, and lead to the formation of defects or clusters, thereby creating a range of local magnetic fields and environments. This variation manifests as a broadened, asymmetric, or more complex EPR spectrum<sup>73</sup>. The second system is associated with the  $\text{V}_\text{C}$  defect centers, for which no changes in line position or width were observed. The weight parameter shows the direct ratio between the two signals ( $\text{Cu}^{2+}/\text{V}_\text{C}$ ), being normalized with the line width and intensity of the two simulated signals. The  $\text{Cu}^{2+}/\text{V}_\text{C}$  ratio was plotted as a function of the Cu dopant concentration, and is depicted in Figure 3f, where the simulation results are also shown in the inset. The ratio between the two paramagnetic centers increases rapidly until a 0.8% Cu concentration is observed after a slight saturation. These findings are further discussed in the electrochemical section of the manuscript, where they are related to the performance of the designed supercapacitor devices based on the g- $\text{C}_3\text{N}_4$ -ZnO:Cu composites.

The electrochemical measurements of the prepared composites were conducted using them as electrode materials in a symmetric supercapacitor device, with 6 M KOH as an electrolyte and glass fiber as a separator. CV is a powerful technique for electrochemical analysis of supercapacitor devices, as it can provide valuable information on their reaction kinetics, electrical conductivity, and charge storage mechanism.

Figure 4a shows the CV curves for all the tested supercapacitor devices measured at 100 mV/s, while Figure S-CV<sup>†</sup> shows the CV curves of the tested supercapacitors at different scan rates. In Figure 4a all supercapacitor devices display a similar quasi-rectangular shape, slightly deviating from the ideal electric double layer capacitance behavior. An ideal EDLC CV curve is a perfect rectangle, with the data following a constant-current, constant-voltage relationship<sup>74</sup>. However, deviations from this ideal rectangular shape are associated with pseudocapacitance, as faradaic reactions at the electrode/electrolyte interface result in a sharp increase in current at the maximum voltage, as observed in all devices. The rounding of the edges of the perfect rectangle is due to the device's internal resistance, which is common in practical devices<sup>75,76</sup>. The change in CV shape with increasing scan rate evident in all the devices as shown in Figure S-CV<sup>†</sup> and for g- $\text{C}_3\text{N}_4$ -ZnO:Cu0.8 in the inset of Figure 4b, show that at lower scan rate ions have more time to penetrate pores and interact with redox-active sites and the CV curves shifts more to ideal

EDLC shape<sup>77,78</sup>.

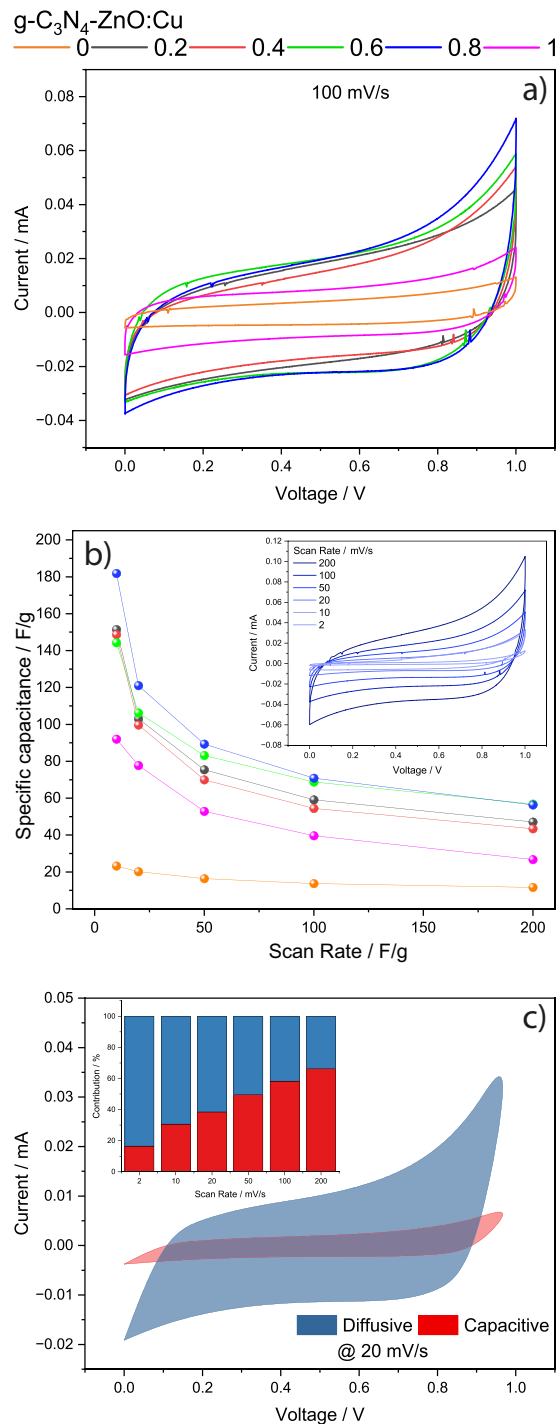


Fig. 4 (a) Cyclic Voltammetry curves for all the tested supercapacitors based on g- $\text{C}_3\text{N}_4$ -ZnO:Cu composites measured at 100 mV/s, (b) specific capacitance values of all supercapacitor devices as a function of the scan rate, with the inset showing the CV curves of the g- $\text{C}_3\text{N}_4$ -ZnO:Cu0.8 devices at different scan rates, and (c) Dunn analysis results showing the capacitive and diffusive contributions to the energy storage mechanism of the supercapacitor device based on g- $\text{C}_3\text{N}_4$ -ZnO:Cu0.8 at a 20 mV/s scan rate, with the inset showing the contributions at different scan rates.

The CV shape suggests that charge storage in all devices is due to the combination of EDLC and pseudocapacitance mech-



anisms. This combination of charge storage mechanisms was expected, as the electrode material possesses EDLC-dominated contributions due to the g-C<sub>3</sub>N<sub>4</sub> carbon-based material, and pseudocapacitance-dominated contributions due to ZnO:Cu<sup>79</sup>. The lowest current response for g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0 compared to other devices suggests the positive impact of Cu ions on the device's electrochemical performance. The trend of the CV results in Figure 4a indicates that there is an increase in current value with increasing Cu doping, with the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8-based supercapacitor device showing the highest current response, and the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu1-based device showing a sudden drop in the current response.

The results indicate that an increase in Cu concentration facilitates charge transport, as evidenced by EPR and XPS. The Cu-doped ZnO is primarily responsible for the pseudocapacitance mechanism in the supercapacitor device, as metal oxides can undergo fast, reversible redox reactions and ion intercalation<sup>80</sup>. These results suggest that increased Cu doping increases the number of Cu<sup>2+</sup> centers, as shown by EPR, thereby further facilitating the redox reaction. However, once a certain dopant threshold is reached, any further increase in dopant concentration results in a sudden drop in current response, which is also visible in Figure 3f, which shows that the Cu<sup>2+</sup>/V<sub>C</sub> ratio starts saturating as the concentration increases above 0.8%. The specific capacitance is a key performance parameter for supercapacitors, providing information on their energy and power density. The specific capacitance can be calculated from the CV curve at different scan rates using the following equation:

$$C_{sp} = \frac{A}{2mv\Delta V} \quad (8)$$

where  $C_{sp}$  is the specific capacitance,  $A$  is the area enclosed by the CV curve,  $m$  is the mass of the active electrode material,  $v$  is the scan rate at which the CV curves were acquired, and  $\Delta V$  is the potential window.

Figure 4b shows the specific capacitance values of all the designed supercapacitor devices plotted as a function of the scan rate, in line with the previous observations, g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8 shows the highest specific capacitance value at each scan rate closely followed by the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.6-based supercapacitor device, but the graph shows that at lower scan rates 0.2, 0.4 and 0.6% Cu-based device show almost identical specific capacitance values. The g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0-based device shows the lowest specific capacitance value while the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu1-based device shows second lowest, which further validates our previous observation that although the Cu doping plays a critical role in improving the performance, after a certain doping percentage, increase in Cu content tends to limit the charge transport, probably due to structural degradation, which is affecting the surface area of the material and decreasing the capacitance value by blocking potential electrochemical sites. As evidenced in Figure 5, where the specific capacitance of the supercapacitor devices based on the composite materials is plotted as a function of the Cu<sup>2+</sup>/V<sub>C</sub> ratio, shows that this is a key parameter that needs fine tuning to enhance the energy storage properties of the materials, playing a key role in the charge transfer and storage processes. An optimum around 0.8%

Cu doping concentration is observable.

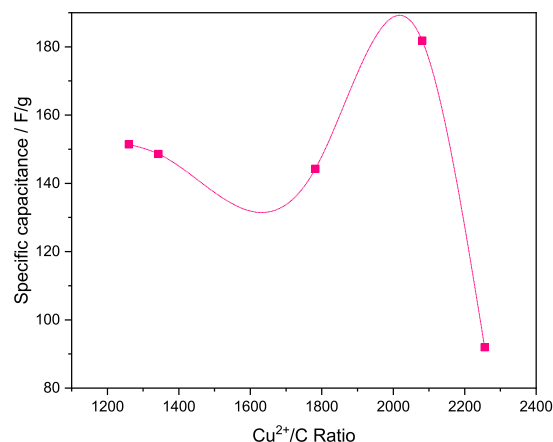


Fig. 5 Specific capacitance values as a function of the Cu<sup>2+</sup>/V<sub>C</sub> ratio, obtained from the simulated EPR spectra.

To analyse the energy storage mechanism in the best-working composite, the contributions of diffusive and capacitive energy storage were assessed using the Dunn method<sup>81</sup>. The equation used to calculate the contributions is expressed as

$$I(v) = k_1 v + k_2 \sqrt{v} \quad (9)$$

where  $k_1$  represents the capacitive current,  $k_2$  represents the diffusive current, and  $v$  is the scan rate. The relative contributions of each mechanism can be quantified by analyzing the slope and intercept of a Dunn plot. Figure 4c shows the CV representation of the diffusive and capacitive contributions at 20 mV/s, with the inset presenting these values at different scan rates. The Dunn analysis complements the observation of the CV results well, as it also indicates that the charge storage used both capacitive and diffusive mechanisms at each scan rate. The Dunn analysis, together with the CV curves shown in the inset of Figure 4b, reveals a gradual transition in the charge-storage mechanism from EDLC dominance at high scan rates to increased surface-confined pseudocapacitance at lower scan rates. As the scan rate decreases, the diffusive contribution extracted from the Dunn analysis becomes more significant, indicating that ion transport within the pores and active faradaic sites increasingly contributes to charge storage. Correspondingly, the CV profiles become more rectangular at lower scan rates because ions have sufficient time to penetrate the pores and access redox-active sites. This enhanced ionic accessibility allows the faradaic reactions to proceed more completely and reversibly, resulting in smoother, more ideal capacitive behavior and demonstrating that pseudocapacitive processes dominate the overall charge storage at low scan rates<sup>77,78</sup>.

A Nyquist plot is a common representation of electrochemical impedance spectroscopy (EIS) data, where the imaginary impedance component is plotted as a function of the real component. Such plots provide information about electrode conductivity, electrolyte resistance, and double-layer capacitance. Figure 6a presents the Nyquist plots of all tested supercapacitor devices. Each device exhibits an incomplete semicircle; the lowest-performing device, g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu1, displays the largest semicir-



cle diameter, corresponding to the highest impedance. In contrast, the other devices show nearly similar semicircle diameters. However, the best-performing devices, g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8 and g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.6, possess the lowest real impedance values<sup>82,83</sup>.

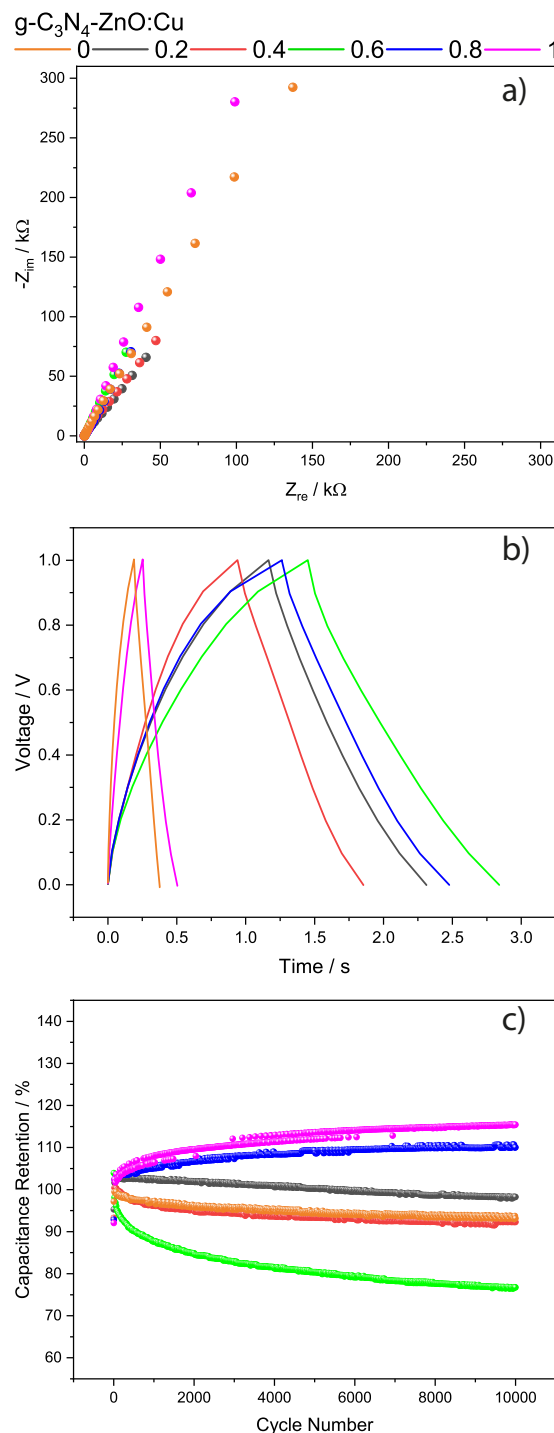


Fig. 6 (a) Nyquist plot, (b) galvanostatic charge-discharge measurements, and (c) capacitance retention as a function of the cycle number of the composite-based supercapacitor devices.

To extract quantitative parameters from the Nyquist plots, equivalent circuit fitting (Z-Fit) was performed. The fitted curves and the corresponding equivalent circuits are shown in Figure S-

PEIS<sup>†</sup>. In the circuit,  $R_1$  ( $R_s$ ) represents the solution resistance, corresponding to the intrinsic resistance between the electrode and electrolyte, while  $R_2$  ( $R_{ct}$ ) denotes the charge-transfer resistance.  $Q$  refers to a constant phase element (CPE), which accounts for deviations from ideal capacitor behavior. Specifically,  $Q_1$  corresponds to the double-layer capacitance ( $C_{dl}$ ), and  $C_2$  or  $Q_2$  represents the pseudocapacitance, as discussed in the CV section. The element W corresponds to Warburg impedance, arising from ion diffusion processes in the electrolyte at low frequencies<sup>84</sup>. The extracted parameter values from the equivalent circuit fitting are summarized in Table 3. Low  $R_s$  and  $R_{ct}$  values, along with a high  $C_{dl}$  value for the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8-based SC design, highlight the superior performance of this device.

Figure 6b presents the galvanostatic charge-discharge measurements of the composite-based supercapacitor devices, carried out at a current density of 0.8 A/g within a voltage window of 0 - 1 V. The charge-discharge curves reveal the pseudocapacitance contribution, consistent with the CV and Dunn analysis results. In an ideal EDLC-type device, the GCD profile exhibits a perfectly triangular shape with complete symmetry between the charge and discharge curves<sup>85,86</sup>. However, the presence of pseudocapacitance distorts this ideal shape<sup>85</sup>. As shown in Figure 6b, a plateau region appears at the maximum charging voltage, indicating the occurrence of faradaic processes<sup>85</sup>. The g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8 and g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.6-based SC devices, which also delivered the highest specific capacitance values, exhibit the longest discharge times. In contrast, the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu1-based device, with the lowest specific capacitance, shows the shortest discharge time, demonstrating consistency across different measurements. S-GCD<sup>†</sup> shows the charge-discharge curves measured at various current densities to analyze device performance further. The graphs follow a similar trend of discharge time as observed in Figure 6b, indicating that with the increasing current density, the device's performance remains stable. Figure 6c illustrates the cyclic stability of all devices, evaluated over 10,000 charge-discharge cycles by plotting the capacitance retention as a function of the cycle number. The results indicate good cyclic stability across all devices, with the g-C<sub>3</sub>N<sub>4</sub>-ZnO: Cu0.6 device showing the lowest retention, maintaining nearly 80% of its initial capacitance after 10,000 cycles. In contrast, the g-C<sub>3</sub>N<sub>4</sub>-ZnO: Cu0.8-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu1 and based supercapacitor demonstrates outstanding stability, with capacitance retention gradually increasing during cycling and reaching almost 110% after 10,000 cycles. This enhancement in retention can be attributed to the progressive activation of electrode materials, improved electrode wettability, and/or structural modifications that provide more accessible active sites for ion storage. Figure S-Ret<sup>†</sup> shows the first and last five charge-discharge cycles for C<sub>3</sub>N<sub>4</sub>-ZnO: Cu0.8 and C<sub>3</sub>N<sub>4</sub>-ZnO: Cu1 based devices, which showed retention higher than 100% to ensure the charge phenomenon is intact and there is no noise or artifact in the signal.

Table 3 shows the energy and power density values for all the supercapacitor devices tested in this study, derived from the highest specific capacitance value. The energy and power density were calculated using the following equations:





Table 3 Electrochemical parameters of the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu-based supercapacitor devices extracted from the equivalent circuit fitting along with the calculated percent error (PE).

Sample	$R_s$ / $\Omega$	$R_{ct}$ / $\Omega$	$C_{dl}$ / $\mu\text{F}$	$W_2$ / $\Omega/\sqrt{s}$	$C_{sp}$ / $\text{F/g}$	$E_D$ / $\text{Wh/kg}$	$P_D$ / $\text{W/kg}$
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0	4.587 (3.2%)	190 (4%)	20.4 (3.6%)	513 (5%)	23.21	3.22	118
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.2	0.345 (4.3%)	115 (3%)	40.00 (4%)	243 (4.2%)	151.4	21.02	764.00
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.4	0.2721 (3.2%)	695 (2.8%)	45.14 (6%)	236 (4.6%)	148.6	20.63	750.18
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.6	0.255 (7.2%)	885 (6.3%)	155.00 (2.2%)	207 (3.7%)	144.25	20.03	739.63
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.8	0.2989 (1.8%)	177 (4.6%)	204.00 (3.3%)	167 (4.2%)	181.83	25.24	917.81
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu1	0.778 (2.4%)	367 (4.7%)	30.13 (3.6%)	396 (3.7%)	91.99	12.77	464.36

$$E_D = \frac{0.5C_{sp}\Delta V}{3.6} \quad (10)$$

$$P_D = \frac{E_D}{\Delta t} 3600 \quad (11)$$

where  $E_D$  and  $P_D$  are the energy and power density, respectively,  $C_{sp}$  is the specific capacitance,  $\Delta V$  is the potential window, and  $\Delta t$  is the discharge time. Energy and power density are the standard comparison metrics for energy storage devices, and the best-performing g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8-based device showed the highest energy and power density values of 25.24 W/kg and 917.81 Wh/kg, respectively. Table 4 summarizes the state-of-the-art performance of previously reported g-C<sub>3</sub>N<sub>4</sub>- and metal-oxide-based supercapacitor systems. Although a direct one-to-one comparison with our device is challenging, as there are various parameters in superapacitor testing, like different electrolytes (acidic/neutral/alkaline), different device configurations (three-electrode vs. two-electrode), and different mass-loadings and calculation approaches, and since our measurements were performed in a two-electrode configuration using 6 M KOH electrolyte and without any conductive additive such as carbon, so, it would be highly unlikely to get a perfect match of parameters for direct comparison.

Material	$C_{ps}$ / $\text{F/g}$	$E_D$ / $\text{Wh/kg}$	$P_D$ / $\text{W/kg}$	Ref.
ZnO	86.4	1.8	2.9	87
ZnO:Cu1%	86.6	48	75	88
g-C <sub>3</sub> N <sub>4</sub> -ZnO	408	-	-	36
g-C <sub>3</sub> N <sub>4</sub> -CuO	98	14.8	-	89
g-C <sub>3</sub> N <sub>4</sub> /TiO <sub>2</sub> /PP	1216	65.6	249.9	90
g-C <sub>3</sub> N <sub>4</sub> /NiO	127	11	600	91
g-C <sub>3</sub> N <sub>4</sub> /ZnO	311	97.43	750	92
g-C <sub>3</sub> N <sub>4</sub> /Fe <sub>2</sub> O <sub>3</sub> :Cu	244	5.31	1000	93
g-C <sub>3</sub> N <sub>4</sub>	78	19.33	500	94
g-C <sub>3</sub> N <sub>4</sub> -ZnO:Cu0.8	181.83	25.24	917.81	t.w.

Table 4  $C_{sp}$ ,  $E_D$ , and  $P_D$  values of supercapacitor devices using electrode materials based on g-C<sub>3</sub>N<sub>4</sub>/Metal oxide composites already reported in the literature. Legend PP - polypyrrole.

In this regard, Table 4 provides a general overview of the performance of these electrode materials, which helps contextualize the performance of our tested device. The best-performing electrode in our study, g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8, exhibited a significantly higher specific capacitance than pristine g-C<sub>3</sub>N<sub>4</sub> and Cu-

doped ZnO electrodes reported in the literature, underscoring the promising electrochemical properties of the g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu composite. Furthermore, several studies listed in Table 4 report g-C<sub>3</sub>N<sub>4</sub>-metal oxide composites with capacitance values comparable to or lower than ours; in contrast, our device demonstrates competitive or superior specific capacitance, energy density, and power density. These results highlight the excellent intrinsic electrochemical performance and strong potential of g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu composites as advanced electrode materials for supercapacitors.

## 4 Conclusions

The g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu nanocomposites were systematically investigated as electrode materials for supercapacitors. The nanocomposite samples were synthesized in several stages: In the first stage, separate components were formed, particularly g-C<sub>3</sub>N<sub>4</sub> via thermal decomposition of urea and Cu-doped ZnO nanoflowers via chemical precipitation. These components were connected using polyallylamine hydrochloride. SEM and TEM investigations revealed uniformly distributed ZnO:Cu nanoflowers embedded within g-C<sub>3</sub>N<sub>4</sub> layers. The incorporation and oxidation state of copper were confirmed using XPS and EPR spectroscopy. Doping introduces several lattice defects in the ZnO lattice, either via substitution or interstitial incorporation, thereby affecting its structural and optical properties. Thus, the developed composite samples have a crystalline structure with particle sizes ranging from 11 to 14 nm, and the band gap value decreases from 3.15 to 3.08 eV. Electrochemical analysis confirmed that the charge storage mechanism originates from a synergistic contribution of EDLC and pseudocapacitance, with Cu doping enhancing conductivity and active site availability. The g-C<sub>3</sub>N<sub>4</sub>-ZnO:Cu0.8-based supercapacitor device exhibited the best overall performance, showing the highest specific capacitance (181.8 F/g), energy density (25.24 Wh/kg), and power density (917.81 W/kg). At the same time, it demonstrated excellent long-term stability, with 110% capacitance retention after 10,000 cycles. The importance of the ratio of carbon-based defect centers in the carbonaceous material g-C<sub>3</sub>N<sub>4</sub> to the concentration of Cu<sup>2+</sup> ions in ZnO was demonstrated, underscoring the crucial role of defect engineering in the design of supercapacitor electrode materials. These results highlight the potential of Cu-doped ZnO when used in composite materials with carbon-based materials, such as g-C<sub>3</sub>N<sub>4</sub>, making them promising candidates for next-generation energy storage devices.



## Author Contributions

**Ana Varadi:** Formal analysis, Investigation, Writing - Original Draft, Writing - Review & Editing; **Anca Silvestru:** Formal analysis, Investigation, Writing - Review & Editing; **Adriana Popa:** Formal analysis, Investigation, Writing - Original Draft, Writing - Review & Editing; **Dana Toloman:** Formal analysis, Investigation, Writing - Original Draft, Writing - Review & Editing; **Arpad Mihai Rostas:** Formal analysis, Writing - Original Draft, Writing - Review & Editing, Visualization; **Ameen Uddin Ammar:** Formal analysis, Writing - Original Draft, Writing - Review & Editing; **Ion Nesterovschi:** Investigation, Writing - Review & Editing; **Maria Mihet:** Investigation, Writing - Review & Editing; **Cristian Leostean:** Formal analysis, Investigation, Writing - Review & Editing; **Sergiu Macavei:** Formal analysis, Investigation, Writing - Review & Editing; **Lucian Barbu-Tudoran:** Formal analysis, Investigation, Writing - Review & Editing; **Maria Stefan:** Conceptualization, Validation, Formal analysis, Investigation, Writing - Original Draft, Writing - Review & Editing, Visualization, Supervision.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 N. I. Jalal, R. I. Ibrahim and M. K. Oudah, *Journal of Physics: Conference Series*, 2021, p. 012015.
- 2 V. Joshi, V. Bachhar, S. S. Mishra, R. K. Shukla and M. Duseja, *Journal of Molecular Structure*, 2025, 143550.
- 3 C. Yang, D. Zhang, D. Wang, H. Luan, X. Chen and W. Yan, *ACS applied materials & interfaces*, 2023, **15**, 5811–5821.
- 4 A. Berrueta, A. Ursua, I. San Martin, A. Eftekhari and P. Sanchis, *Ieee Access*, 2019, **7**, 50869–50896.
- 5 J. Guo, D. Zhang, T. Li, J. Zhang and L. Yu, *Journal of Colloid and Interface Science*, 2022, **606**, 261–271.
- 6 H. Aslam, A. Umar, M. U. Khan, T. Trivedi, G. Ezhilarasan, D. Bhanot, N. Abbas and K. F. Fawy, *Reviews in Inorganic Chemistry*, 2025.
- 7 W. Yang, M. Ni, X. Ren, Y. Tian, N. Li, Y. Su and X. Zhang, *Current Opinion in Colloid & Interface Science*, 2015, **20**, 416–428.
- 8 M. Saranya, R. Ramachandran and F. Wang, *Journal of Science: Advanced Materials and Devices*, 2016, **1**, 454–460.
- 9 D. P. Chatterjee and A. K. Nandi, *Journal of Materials Chemistry A*, 2021, **9**, 15880–15918.
- 10 V. V. Obreja, *Physica E: Low-dimensional Systems and Nanostructures*, 2008, **40**, 2596–2605.
- 11 T. Jasna, S. K. Subrahmanian, V. Kavya, A. Haritha,

- T. Karthika, M. Shemeena, P. Jabira and B. N. Narayanan, *Journal of Physics D: Applied Physics*, 2024, **58**, 035504.
- 12 T. Tene, S. Bellucci, M. Guevara, P. Romero, A. Guapi, L. Gahramanli, S. Straface, L. S. Caputi and C. Vaca-cela Gomez, *Batteries*, 2024, **10**, 256.
- 13 B. Zhao, P. Liu, Y. Jiang, D. Pan, H. Tao, J. Song, T. Fang and W. Xu, *Journal of power sources*, 2012, **198**, 423–427.
- 14 Y. Wang, K. Xue, X. Zhang, X. Zhang, P. Ma, B. Yang, S. Xu and J. Lang, *Electrochimica Acta*, 2023, **441**, 141829.
- 15 I. Shaheen, I. Hussain, T. Zahra, M. S. Javed, S. S. A. Shah, K. Khan, M. B. Hanif, M. A. Assiri, Z. Said, W. U. Arifeen *et al.*, *Journal of Energy Storage*, 2023, **72**, 108719.
- 16 H. Zhang, D. Zhang, R. Mao, L. Zhou, C. Yang, Y. Wu, Y. Liu and Y. Ji, *Nano Energy*, 2024, **127**, 109753.
- 17 J. Miao and B. Liu, in *Semiconductor Nanowires*, Elsevier, 2015, pp. 3–28.
- 18 F. Pogacean, M. Ștefan, D. Toloman, A. Popa, C. Leostean, A. Turza, M. Coros, O. Pana and S. Pruneanu, *Nanomaterials*, 2020, **10**, 1473.
- 19 I. Ayoub, V. Kumar, R. Abolhassani, R. Sehgal, V. Sharma, R. Sehgal, H. C. Swart and Y. K. Mishra, *Nanotechnology Reviews*, 2022, **11**, 575–619.
- 20 D. Toloman, A. Gungor, A. Popa, M. Stefan, S. Macavei, L. Barbu-Tudoran, A. Varadi, I. D. Yildirim, R. Suci, I. Nesterovschi *et al.*, *Ceramics International*, 2025, **51**, 353–365.
- 21 S. Sivakumar, Y. Robinson and N. A. Mala, *Applied Surface Science Advances*, 2022, **12**, 100344.
- 22 M. R. Pallavolu, J. Nallapureddy, R. R. Nallapureddy, G. Nee-lima, A. K. Yedluri, T. K. Mandal, B. Pejja and S. W. Joo, *Journal of Alloys and Compounds*, 2021, **886**, 161234.
- 23 S. Kumar, F. Ahmed, N. M. Shaalan, N. Arshi, S. Dalela and K. H. Chae, *Nanomaterials*, 2023, **13**, 2222.
- 24 A. U. Ammar, M. H. Aleinawi, M. Stefan, A. Gungor, A. Popa, D. Toloman, K. Maškarić, E. Saritas, L. Barbu-Tudoran, S. Macavei *et al.*, *Electrochimica Acta*, 2025, **519**, 145806.
- 25 M. Stefan, D. Toloman, A. Popa, A. Mesaros, O. Vasile, C. Leostean and O. Pana, *Journal of Nanoparticle Research*, 2016, **18**, 59.
- 26 M. Zhi, C. Xiang, J. Li, M. Li and N. Wu, *Nanoscale*, 2013, **5**, 72–88.
- 27 S. Tamang, S. Rai, R. Bhujel, N. K. Bhattacharyya, B. P. Swain and J. Biswas, *Journal of Alloys and Compounds*, 2023, **947**, 169588.
- 28 D. Zhang, C. Jiang and X. Zhou, *Talanta*, 2018, **182**, 324–332.
- 29 G. Yu, X. Xie, L. Pan, Z. Bao and Y. Cui, *Nano Energy*, 2013, **2**, 213–234.
- 30 H. Zhang, D.-Z. Zhang, D.-Y. Wang, Z.-Y. Xu, Y. Yang and B. Zhang, *Rare Metals*, 2022, **41**, 3117–3128.
- 31 Y. Wang, Y. Liu and J. Zhang, *Journal of Nanoparticle Research*, 2015, **17**, 1–10.
- 32 M. Ramezani, M. Fathi and F. Mahboubi, *Electrochimica Acta*, 2015, **174**, 345–355.
- 33 C. Xiang, M. Li, M. Zhi, A. Manivannan and N. Wu, *Journal of Power Sources*, 2013, **226**, 65–70.
- 34 Z. J. Han, S. Pineda, A. T. Murdock, D. H. Seo, K. K. Ostrikov



- and A. Bendavid, *Journal of Materials Chemistry A*, 2017, **5**, 17293–17301.
- 35 B. Wang, D. Guan, Z. Gao, J. Wang, Z. Li, W. Yang and L. Liu, *Materials chemistry and physics*, 2013, **141**, 1–8.
- 36 S. Savarimuthu, M. Selvaraj and H. Chuang, *Inorganic Chemistry Communications*, 2025, **178**, 114555.
- 37 M. Sayed, B. Zhu, P. Kuang, X. Liu, B. Cheng, A. A. A. Ghamdi, S. Wageh, L. Zhang and J. Yu, *Advanced Sustainable Systems*, 2022, **6**, 2100264.
- 38 D. Dvoranová, M. Mazúr, I. Papailias, T. Giannakopoulou, C. Trapalis and V. Brezová, *Catalysts*, 2018, **8**, 47.
- 39 M. Stefan, A. M. Rostas, A. U. Ammar, A. Gungor, E. Saritas, D. Toloman, A. Varadi, S. Macavei, L. Barbu-Tudoran, C. Leostean *et al.*, *Energy & Fuels*, 2024, **38**, 19088–19099.
- 40 A. B. Djurišić, X. Chen, Y. H. Leung and A. M. C. Ng, *Journal of Materials Chemistry*, 2012, **22**, 6526–6535.
- 41 G.-L. Li, S.-Y. Wu, X.-F. Hu, B.-H. Teng and M.-H. Wu, *Journal of Applied Spectroscopy*, 2016, **83**, 374–378.
- 42 *The Journal of Chemical Physics*, 2021, **154**, 124710.
- 43 K. T. Lee, S. Jeong and J. Cho, *Accounts of chemical research*, 2013, **46**, 1161–1170.
- 44 R. DANESI, 2018.
- 45 M. Q. Alfaifi and A. A. Bagabas, *Adv. Mater. Sci.*, 2019, **4**, 1–10.
- 46 X. Shao, D. Zhang, M. Tang, H. Zhang, Z. Wang, P. Jia and J. Zhai, *Chemical Engineering Journal*, 2024, **495**, 153676.
- 47 M. M. Khan, S. Kumar, A. N. Alhazaa and M. Al-Gawati, *Materials Science in Semiconductor Processing*, 2018, **87**, 134–141.
- 48 X. Guo, J. Duan, C. Li, Z. Zhang and W. Wang, *Journal of Materials Science*, 2020, **55**, 2018–2031.
- 49 M. Sher, M. Javed, S. Shahid, S. Iqbal, M. A. Qamar, A. Bahadur and M. A. Qayyum, *RSC advances*, 2021, **11**, 2025–2039.
- 50 Y. Wang, R. Shi, J. Lin and Y. Zhu, *Energy & Environmental Science*, 2011, **4**, 2922–2929.
- 51 D. Sudha, E. R. Kumar, S. Shanjitha, A. M. Munshi, G. A. Al-Hazmi, N. M. El-Metwaly and S. J. Kirubavathy, *Ceramics International*, 2023, **49**, 7284–7288.
- 52 V. Mursyalaat, V. Variiani, W. Arsyad and M. Firihi, *Journal of Physics: Conference Series*, 2023, p. 012042.
- 53 D. Toloman, A. Popa, M. Stefan, O. Pana, T. D. Silipas, S. Macavei and L. Barbu-Tudoran, *Materials Science in Semiconductor Processing*, 2017, **71**, 61–68.
- 54 B. Chander Joshi and A. K. Chaudhri, *ACS omega*, 2022, **7**, 21877–21881.
- 55 M. Masar, H. Ali, A. C. Guler, M. Urbanek, P. Urbanek, B. Hanulikova, H. Pistekova, A. Annusova, M. Machovsky and I. Kuritka, *Colloids and Surfaces A: Physicochemical and Engineering Aspects*, 2023, **656**, 130447.
- 56 D. Toloman, A. Popa, M. Stefan, T. D. Silipas, R. C. Suciui, L. Barbu-Tudoran and O. Pana, *Optical Materials*, 2020, **110**, 110472.
- 57 C. Xia, F. Wang and C. Hu, *Journal of alloys and compounds*, 2014, **589**, 604–608.
- 58 Z. Ma, F. Ren, X. Ming, Y. Long and A. A. Volinsky, *Materials*, 2019, **12**, 196.
- 59 I. Idrees, A. Razzaq, M. Zafar, A. Umer, F. Mustafa, F. Rehman and W. Y. Kim, *Arabian Journal of Chemistry*, 2024, **17**, 105615.
- 60 A. Yavuz, D. Aydin, B. Disli, T. Ozturk, B. Gul, I. H. Gubbuk and M. Ersoz, *Environmental Science and Pollution Research*, 2024, **31**, 44828–44847.
- 61 Q. Xiang, J. Yu and M. Jaroniec, *The Journal of Physical Chemistry C*, 2011, **115**, 7355–7363.
- 62 H. Wang, X. Zhang, J. Xie, J. Zhang, P. Ma, B. Pan and Y. Xie, *Nanoscale*, 2015, **7**, 5152–5156.
- 63 R. Mekkat, E. M. Kutorglo, M. Setka, A. Prokhorov and M. Šošč, *Journal of Photochemistry and Photobiology A: Chemistry*, 2025, 116454.
- 64 M. Sakkaki and S. M. Arab, *Synthesis and Sintering*, 2022, **2**, 176–180.
- 65 T. Suter, V. Brázdová, K. McColl, T. S. Miller, H. Nagashima, E. Salvadori, A. Sella, C. A. Howard, C. W. Kay, F. Corà *et al.*, *The Journal of Physical Chemistry C*, 2018, **122**, 25183–25194.
- 66 D. S. Patrick, A. Govind, P. Bharathi, M. K. Mohan, S. Harish, J. Archana and M. Navaneethan, *Applied Surface Science*, 2023, **609**, 155337.
- 67 D. Toloman, A. Popa, M. Stan, M. Stefan, G. Vlad, S. Ulinici, G. Baisan, T. D. Silipas, S. Macavei, C. Leostean *et al.*, *Journal of Alloys and Compounds*, 2021, **866**, 159010.
- 68 A. U. Ammar, I. D. Yildirim, M. H. Aleinawi, M. Buldu-Akturk, N. S. Turhan, S. Nadupalli, A. M. Rostas and E. Erdem, *Materials Research Bulletin*, 2023, **160**, 112117.
- 69 P. Xia, B. Cheng, J. Jiang and H. Tang, *Applied Surface Science*, 2019, **487**, 335–342.
- 70 B. Babu, T. Aswani, G. T. Rao, R. J. Stella, B. Jayaraja and R. Ravikumar, *Journal of magnetism and magnetic materials*, 2014, **355**, 76–80.
- 71 D. R. Lorenz, J. R. Wasson, P. R. Johnson and D. A. Thornton, *Journal of Inorganic and Nuclear Chemistry*, 1975, **37**, 2297–2302.
- 72 S. Stoll and A. Schweiger, *Journal of magnetic resonance*, 2006, **178**, 42–55.
- 73 W. Hagen, in *Practical approaches to biological inorganic chemistry*, Elsevier, 2020, pp. 121–154.
- 74 Y. Gogotsi and R. M. Penner, *Energy storage in nanomaterials—capacitive, pseudocapacitive, or battery-like?*, 2018.
- 75 J. P. A. dos Santos, F. C. Rufino, J. I. Y. Ota, R. C. Fernandes, R. Vicentini, C. J. Pagan, L. M. Da Silva and H. Zanin, *Journal of Energy Chemistry*, 2023, **80**, 265–283.
- 76 W. K. Chee, H. N. Lim, I. Harrison, K. F. Chong, Z. Zainal, C. H. Ng and N. M. Huang, *Electrochimica Acta*, 2015, **157**, 88–94.
- 77 N. G. Hörmann and K. Reuter, *Journal of Physics: Condensed Matter*, 2021, **33**, 264004.
- 78 H. Zhu, Y. Li, Y. Song, G. Zhao, W. Wu, S. Zhou, D. Wang and W. Xiao, *Materials technology*, 2020, **35**, 135–140.
- 79 D. A. Bogachev, Y. M. Volfkovich and S. Martemianov, *Journal*



- of *Electroanalytical Chemistry*, 2023, **935**, 117322.
- 80 M. Alexandreli, C. B. Brocchi, D. M. Soares, W. G. Nunes, B. G. Freitas, F. E. de Oliveira, L. E. C. A. Schiavo, A. C. Peterlevitz, L. M. da Silva and H. Zanin, *Journal of Energy Storage*, 2021, **42**, 103052.
  - 81 W. Pholauyphon, P. Charoen-Amornkitt, T. Suzuki and S. Tsushima, *Electrochemistry Communications*, 2024, **159**, 107654.
  - 82 P. Widiatmoko, N. Nugraha, H. Devianto, I. Nurdin and T. Prakoso, IOP Conference Series: Materials Science and Engineering, 2020, p. 012040.
  - 83 A. C. Lazanas and M. I. Prodromidis, *ACS measurement science au*, 2023, **3**, 162–193.
  - 84 M. Y. Perdana, B. A. Johan, M. Abdallah, M. E. Hossain, M. A. Aziz, T. N. Baroud and Q. A. Drmash, *The Chemical Record*, 2024, **24**, e202400007.
  - 85 Y. Shao, M. F. El-Kady, J. Sun, Y. Li, Q. Zhang, M. Zhu, H. Wang, B. Dunn and R. B. Kaner, *Chemical reviews*, 2018, **118**, 9233–9280.
  - 86 A. Singh and A. Chandra, *Enhancing specific energy and power in asymmetric supercapacitors-a synergetic strategy based on the use of redox additive electrolytes*, *Sci. Rep.* **6** (2016) 25793, 2016.
  - 87 I. Shaheen, K. S. Ahmad, C. Zequine, R. K. Gupta, A. G. Thomas and M. Azad Malik, *Environmental Technology*, 2022, **43**, 605–616.
  - 88 A. U. Ammar, F. Bakan-Misirlioglu, M. H. Aleinawi, G. Franzo, G. G. Condorelli, F. N. T. Yesilbag, Y. O. Yesilbag, S. Mirabella and E. Erdem, *Materials Research Bulletin*, 2023, **165**, 112334.
  - 89 S. Surender, M. Kavipriyah and S. Balakumar, *Inorganic Chemistry Communications*, 2023, **150**, 110557.
  - 90 S. Ahlawat and S. Lata, *Materials Research Bulletin*, 2023, **165**, 112328.
  - 91 L. M. Jose, N. S. George, S. A. Kadam, D. Nayana and A. Aravind, *Inorganic Chemistry Communications*, 2025, **178**, 114527.
  - 92 A. Kumar, H. Mittal and M. Khanuja, *Journal of Energy Storage*, 2023, **67**, 107496.
  - 93 C. Harak, V. Kadam, R. Gavhane, S. Balgude, A. Rakshe, N. Brahmanekar, S. Uke, D. Satpute, H. Pawar and S. Mardikar, *RSC advances*, 2024, **14**, 4917–4929.
  - 94 L. G. Ghanem, M. A. Hamza, M. M. Taha and N. K. Allam, *Journal of Energy Storage*, 2022, **52**, 104850.





“All information supporting the findings of this study are fully included within the main article and its accompanying Supplementary Information files. No additional datasets were generated or analyzed beyond those provided.”

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