

Green synthesis of CuO nanoparticles using Jasmin sambac extract: Conditions optimization and photocatalytic degradation of Methylene Blue dye 2

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Abstract

This investigation utilized an eco-friendly method to produce copper oxide nanoparticles (CuO NPs) through the utilization of *Jasminium sambac* leaf extract. Optimization of pH, temperature, reaction time, and copper sulfate concentration were performed. Characterization via UV-Vis, SEM, XRD, FTIR, and EDX techniques revealed a peak at 243 nm in UV-Vis, and an FTIR band at 590 cm⁻¹ confirmed Cu(II) ion reduction to CuO NPs. pH, temperature, reaction time, and copper sulfate concentration were identified as crucial synthesis parameters. EDX analysis indicated a composition of 75.94% copper and 24.06% oxygen. CuO NPs were dispersive, monoclinic, crystalline, and pure, with an average particle size of 13.4 nm. Photocatalytic experiments achieved 97% Methylene Blue dye degradation in 210 minutes, highlighting the efficacy of the green synthesis method for CuO NPs with significant photocatalytic activity in wastewater remediation.

Keywords: Green synthesis; CuO nanoparticles; Photocatalysis; Dye degradation

1. Introduction

Nanotechnology's impact in recent decades has been significant, especially due to the unique properties of nanoparticles arising from their nanoscale size. With a higher surface-to-volume ratio, nanoparticles find versatile applications across various fields (Alfanaar et al., 2021, Nagesh et al., 2022, Sivakavinesan et al., 2022). In the NP synthesis, a reduction of metal ions is performed using an appropriate reducing agent. In view of the existing situation of environmental contamination (Iqbal et al., 2019), The demand for eco-friendly nanomaterial preparation has grown. Conventional synthesis methods involve hazardous reagents, extended reaction times, and high costs. Green chemistry tackles these challenges by employing economical and environmentally friendly agents. Plant extracts, rich in natural carboxyl and hydroxyl groups, present ideal substitutes for chemicals in environmentally friendly synthesis.

They adeptly reduce metal ions to metal atoms at the nano-scale, serving as dual-function agents for both reduction and stabilization (Shammout and Awwad 2021). The *Jasminium sambac* leaves are rich in phenolic compounds and can be used for green synthesis of NP since it is rich in antioxidant activity (Widowati et al., 2018). Green synthesis is more cost-effective and time-efficient than traditional physical and chemical methods for nanoparticle synthesis, requiring no normal energy, pressure, or heating conditions, and avoiding the use of harmful chemicals (Naseer et al., 2020, Amer and Awwad 2021, Shammout and Awwad 2021).

The textile industry contributes significantly to water pollution through the discharge of industrial waste containing colored chemicals. This leads to reduced dissolved oxygen, increased turbidity, and interference with natural photosynthesis in water bodies. The MB dye, commonly used in aquaculture for fish disease treatment and as a mild antifungal agent, poses harm to aquatic life (Iqbal et al., 2019). The environmental impact of MB dye varies based on its application, concentration, and disposal. Caution in use benefits water quality monitoring and medical diagnostics. Improper handling or disposal can result in negative consequences, including water contamination and harm to aquatic life (Ayach et al., 2017, Iqbal et al., 2019). Effective treatment of textile wastewater, laden with toxic dyes harmful to living organisms, is crucial before discharge. Various methods, such as adsorption, chemical coagulation, membrane separation, photocatalytic degradation, and biological treatment, are employed for addressing water with soluble dyes (Iqbal 2016, Djehaf et al., 2017). Because it removes pollutants and toxins from sewage and industrial effluents, wastewater treatment is critical for environmental and public health protection (Yildirim and Sasmaz 2017). There are various wastewater treatment technologies (Physical, chemical and biological), each with its own set of benefits and drawbacks (Khera et al., 2020, Neolaka et al., 2023). On the other hand, photocatalysis is an effective method for the elimination of toxic dyes from industrial effluents,

which destruct the organic pollutants into water, carbon dioxide and inorganic ions (Ali et al., 2021).

The goal of this study was to synthesize CuO NPs from *Jasminium sambac* leaf extract, which is high in flavonoids, phenolics, saponins, and steroids. UV-Vis, SEM, XRD, FTIR, and EDX techniques were used to characterize the samples. This study also examined the influence of process variables on the photocatalytic activity (PCA) in the degradation of Methylene Blue (MB) dye.

2. Experimental

2.1 Reagents and plant samples preparation

Analytical grade chemicals were used in the green synthesis of CuO NPs. Sigma Aldrich provided $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ and Methylene Blue, while Merck provided NaOH. The leaves of *Jasminium sambac* (Motia) were collected in Sialkot, Pakistan. Among various extraction techniques (Kusuma and Mahfud 2017, Kusuma and Mahfud 2018), the hydro distillation is a method of extraction that is commonly used for the separation of bioactive compounds from plant materials, such as herbs, flowers, or citrus peels. The leaves were washed, dried in the sun, over-dried (to a constant weight), and ground to a fine powder. In 100 mL of distilled water, 10 g of powdered leaves were heated at 80 °C with continuous stirring for 15 minutes. The extract was cooled and filtered, and the supernatant was stored at 4 °C before being used to make CuO NPs.

2.3. Green synthesis of CuO NPs

For CuO NPs synthesis, $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ was used as a source of Cu^{+2} ion. 20 mL of 1M $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ was added in 80 mL of plant extract solution, stirred and the color changed to brown suspension, filtered and dried at 60 °C for 2 h. The following different parameters were optimized for the synthesis of CuO NPs. Salt concentration was optimized by varying the salt concentration from 0.1 to 2 M with constant pH (pH 8), temperature (60 °C), and time (30 min).

Optimization of pH was done at different pH (2-8 pH) with constant temperature (60°C) salt concentration (1M), and time (30 min). The temperature effect was studied by varying the temperature from 30-90 °C at constant time (30 min), pH (pH 8), and salt concentration (1M). The effect of reaction time was studied in 30-90 min range. The CuO NPs synthesis mechanism prepared via a green route using *Jasmin sambac* extract is depicted in Fig. 1 (Nazir et al., 2023).

Phytochemicals in *Jasminum sambac* extracts, like alkaloids, flavonoids, tannins, terpenoids, phenolic compounds, saponins, glycosides, and carotenoids, play a vital role in reducing metal ions, contributing to the plant's biological and therapeutic properties. Composition variations stem from factors like plant age, environmental conditions, and extraction methods (Kusuma and Mahfud 2017, Putri et al., 2019). *Jasminum sambac* extract analysis revealed the identification of 54 bioactive components in the leaf extract. The diverse array of compounds exhibits known biological properties (Tomar and Rijhwani 2015). The presence of phytochemicals in *Jasminum sambac* suggests potential health benefits and pharmacological activities, given the well-known diverse biological properties of these compounds (Rattan 2023). The synthesis of CuO NPs involves the reduction of metal ions, facilitated by the abundance of bioactive compounds in *Jasminum sambac*. Fig. 1 illustrates the CuO NPs synthesis mechanism using *Jasminum sambac* extract, following a green route (Nazir et al., 2023). Green synthesis using plant extracts offers advantages like reduced energy consumption, eco-friendly reagents, and scalability, with the mechanism influenced by the bioactive content and chosen extraction technique (Kusuma and Mahfud 2016, Kusuma and Mahfud 2017, Kusuma et al., 2018).

2.5. Characterizations

The UV-Visible spectrophotometer was used to measure the wavelength range of 200-800 nm. XRD (Philips PANalytical powder) was used to characterize green synthesized CuO NPs (CuK

radiation of wavelength 0.15406 nm in 15 scan range of 20°-80°). To study the morphology, synthesized nanoparticles were examined by using SEM (TESCAN LMU Vega – Variable pressure Scanning Electron Microscope). EDX analysis was performed using an EDX TESCAN LMU Vega to identify elements and determine the chemical composition of elements in synthesized materials, and an FT-IR spectrum was used to identify the reducing and capping agents present in leaf extract and NPs. The Nicolet 6700 FTIR spectrophotometer with a wavelength range of 400 cm⁻¹ to 4000 cm⁻¹ was used for this analysis.

2.6. Photocatalytic procedure

CuO NPs were used to photocatalyze the degradation of MB dye, which was observed through color change in an aqueous solution exposed to light. After 30 minutes of stirring in the dark, the CuO NPs and dye mixture reached adsorption-desorption equilibrium before being exposed to light (880.50-890.50 W/m² was the intensity of light). A small aliquot of the solution was taken out after a constant interval of time and centrifuged at 3000 rpm for 10 min, before UV visible analysis. Different parameters such as pH (2 -12 pH) were analyzed for optimal pH study with constant parameters like time (105 min), concentration of Dye (20 ppm), and amount of CuO NPs (50 mg). Optimization study on time (0 -120 min) was performed and analyzed with other constant parameters like concentration of Dye (50 ppm), amount of CuO NPs (20 mg), and pH 8. The concentration of dye (10-50 ppm) was analyzed for the optimization study with constant parameters like time (90 min) amount of CuO NPs (20 mg) and amount of CuO NPs (10-50 mg) were analyzed for the degradation of MB dye with constant dye concentration (50 ppm), time (120 min), and at pH 8. Percentage photocatalytic degradation was determined using Eq. 1. All the experiments were run in triplicate and data thus obtained was averaged.

$$\text{Photocatalytic degradation (\%)} = \frac{(A_0 - A_t)}{A_0} \times 100 \quad (1)$$

Where A₀ and A_t represent the absorbance values before and after irradiation.

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3. Results and discussion

3.1. Characterization of NPs

3.2. Process variables effect on the synthesis of CuO NPs

3.2.1. Effect of salt concentration

Figure 2A represents spectra of CuO NPs at different concentrations of $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ (0.1, 0.5, 1, 1.5, and 2 M). At 1M concentration, maximum absorption was observed at 243 nm. Results showed that the intensity of the absorption decreased with the rise of salt amount from 1M to 2M. This may be attributed to the lower concentration of reducing agent and Cu^{+2} ion in higher concentration. Whereas, the minimum value of absorption at 0.1 M concentration may be attributed to the lower concentration of Cu^{+2} ion in the solution. The observations revealed that a salt concentration of 1M was optimum for the CuO NPs synthesis.

3.2.2. Effect of pH

Figure 2B depicts the pH 2, 4, and 6 effect on the formation of CuO NPs. At alkaline pH (8), an intense spectrum was observed at 243 nm indicating the synthesis of stabilized CuO NPs. Whereas at acidic pH, the reducing and stabilizing agent of plant extract was in their protonated form and therefore, absorption was weak. The results indicate that highly stable, small-sized CuO NPs were synthesized at alkaline pH. Similar results were obtained for NPs synthesis using plant extract, indicating that pH can influence NP formation (Piñero et al., 2017, Kredy 2018).

3.2.3. Effect of temperature

Figure 2C revealed a small shift in absorption peak to longer wavelength with the rise in temperature from 30 °C to 90 °C. This shift is due to the increase in particle size at high temperatures. The SEM image also shows a rise in particle size with temperature. Alike findings have also been reported for the synthesis of NPs using plant extract that the temperature may affect the NPs formation significantly (Kredy 2018).

3.2.4. Effect of Reaction Time

The absorption spectra for CuONP syntheses at 30, 60 and 90 min reaction time are shown in Fig. 3A. The results represent absorption peaks at about 224, 234 and 244 nm, respectively. The small shift in the absorption band at a longer wavelength indicates the increase in particle size. This could be better explained in terms of optical band gap energy, which was determined using Eq. 2.

$$(\alpha h\nu)^n = B(h\nu - E_g) \quad (2)$$

Where B is a constant, $h\nu$ is photon energy in eV, h is Planck's constant, ν is photon frequency, E_g is optical band gap energy in eV, n is an exponent transition, and α is absorption coefficient. The increase in reaction time is responsible for the shift in band gap energy (Fig. 3B). As the particle size decreases to the nanoscale, the energy level decreases and electron becomes more confined in the particles. This increases the band gap energy between the valance and conduction bands. This explains why NPs have a higher band gap energy than comparable large-size materials. (Naseer et al., 2020). Furthermore, the XRD analysis results show that the CuO NPs size increases with reaction time, from 13.4 to 15.7 nm.

3.3. XRD analysis

The CuO NPs formation was confirmed by XRD analysis (Philips PANalytical powder) using $\text{CuK}\alpha$ radiation of wavelength 0.15406 nm in a scan range of 20°-80°. X-ray diffraction pattern show (Fig. 4) showed sharp peaks at 35.71°, 38.93°, 48.98° and 62.07°, indicating the formation of the CuO NPs monoclinic and crystalline in nature. The size (D) of CuO NPs obtained by using applying the Debye-Scherrer formula (Eq. 3) was found 13.4 nm.

$$D = \frac{0.9 \lambda}{\beta \cos \theta} \quad (3)$$

The X-ray diffraction patterns of CuO NPs prepared using *Jasminium sambac* leaves extract are closely related to the XRD pattern of CuO NPs previously reported (Das et al., 2013). Also, (Acharyulu et al., 2014, Shi et al., 2017) reported pure crystallite and well-defined CuO NPs using *C. auriculata* and *P. amarus* leaf extract, respectively.

3.4. EDX and SEM analysis

The elemental composition of CuO NPs was validated using energy dispersive X-ray analysis (EDX). The EDX analysis confirms the presence of Cu and O elements, as shown in Fig. 5, with percentages of 75.94 and 24.06, respectively. CuO NPs exhibit a strong diffraction band peak at 1keV, which is typical of copper absorption due to surface plasmon resonance. SEM was used to examine the morphology of green fabricated CuO NPs, and the results are shown in Fig. 6. The average size of CuO NPs was discovered to be consistent with XRD analysis. Because of the interaction between biomolecules capping the individual particle, the CuO NPs had irregular shapes and aggregated.

3.5. FTIR analysis

FT-IR analysis is employed for the identification of functional groups (Syafaatullah and Mahfud 2021), which is used to identify the bioactives in the leaf extract of *Jasminium sambac*. It is also used to study the effective incorporation of phytochemicals in the synthesis of CuO NPs. FTIR spectrum (Fig. 7A) of CuO NPs and leaf extract was recorded in the 400 to 4000 cm^{-1} range. The band at 2718.11 cm^{-1} is ascribed to C-H vibration. While a band at 1024.72 cm^{-1} and 831.71 cm^{-1} are associated with C-N stretching in primary amine and C-C vibrations in the alkyl group present in plant extract. The bands at 2354.86 cm^{-1} show the alkyne C≡C stretching vibration, which indicates the presence of lipid molecules in the leaf extract. The most significant absorption band detected at 491 cm^{-1} is allocated to metal-oxygen vibration (Shi et al., 2017).

3.6. Photocatalytic activity of CuO NPs

3.6.1. Effect of pH

The impact of solution pH on dye degradation was investigated by varying pH levels from 2 to 12. Experiments were conducted with a fixed dye concentration of 20 ppm and 50 mg of CuO NPs. The degradation efficiency of CuO NPs was found to be 52.63, 61.1, 69.51, and 79.82 (%), at pH 2, 4, 6, 8, and 12, respectively (Table 1). In alkaline pH, a negative charge is induced

on the CuONP surface and enhances the formation of reactive species that effectively degrade the cationic dye. At acidic pH, there were no significant electrostatic attractions between the CuONP surface and MB dye molecules. Therefore, the extent of photocatalytic degradation of MB dye was decreased at acidic pH. It is evident from the results that, an increase in pH from 2 to 8 accelerates photocatalytic degradation, while at pH 12, the photocatalytic degradation efficiency of CuO NPs declined. The reason is attributed to the excess of anions formation at $\text{pH} > 8$ (Acharyulu et al., 2014). Anions compete with dye molecules to adsorb on CuONP surface active sites, hindering adsorption and reducing photocatalytic degradation of MB dye. At pH_{pzc} is the point where nanoparticles surface charge became neutral. CuO nanoparticles have an amphoteric behavior, which means they can be positively or negatively charged depending on the pH of their surroundings. Understanding the nature of the binding active site on the surface of CuO nanoparticles requires an understanding of the point of zero charge (pH_{pzc}). (Noreen et al., 2020). For this, a various solution of CuO nanoparticles were prepared in NaCl solution and then, their initial pH was adjusted using NaOH/HCl solutions and final pH values were estimated after 24 h. The $\text{pH} < \text{pH}_{\text{pzc}}$ favors the adsorption of cationic species, while $\text{pH} > \text{pH}_{\text{pzc}}$ favors anionic species adsorption. The pH data thus obtained was used to measure the change in pH values and a plot was drawn between initial pH and ΔpH ($\text{pH}_i - \text{pH}_f$). The data revealed that the values of pH_{pzc} of CuO nanoparticles was 6.27 (Fig. S1). The pH_{pzc} data revealed that the CuO nanoparticle bear positive charge and negative above the pH_{pzc} . Also, the findings of present investigation support these observations. In basic pH, the dye degradation was higher, which is due to the negative charge on the surface of CuONP surface that enhances the reactive species formation (Liang et al., 2017). The cationic dye was effectively attracted to the CuONP surface, enhancing dye removal through the oxidative process, particularly in basic media.

3.6.2. ¹Effect of CuO NP dose

The dose of CuO NPs has a substantial influence on dye destruction. The photocatalytic degradation of dye was increased by increasing the amount of catalyst. As the concentration of CuO NPs increased from 10 mg to 50 mg, the degradation efficiency of MB dye increased from 11.62% to 80.67% (Table S1). This phenomenon is due to the augmentation of active sites on the catalyst, which results in optimal adsorption of dye molecules onto the catalyst's surface and the generation of additional reactive species for dye degradation.

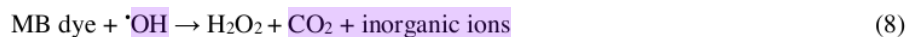
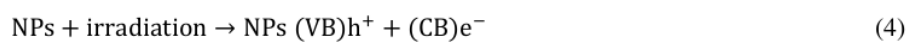
3.6.3. Effect of dye initial concentration

The effect of the concentration of MB dye on photocatalytic degradation was investigated in 10 to 50 mg/L. By enhancing the dye concentration, the dye removal was declined (Table S2). This might be due to the obstruction in the path of incident light due to the increased concentration of dye, which precludes the light from reaching the catalyst surface, thus leading to a decrease in degradation efficiency. Secondly, the adsorption of MB molecules on the catalyst surface suppressed the generation of reactive species, and resultantly, degradation efficiency was decreased.

3.6.4. Effect of contact time

The impact of contact time on photocatalytic degradation is shown in Fig. 7B. The before irradiation of catalysts and dye mixture to the light, the catalysts and dye mixture was stirred in dark for 30 min to check to attain the adsorption-desorption mechanism of dye. Before irradiation to light, 3 mL dye sample was taken and its absorbance was monitored. It was observed that 1.72% dye was adsorbed during the adsorption-desorption equilibrium attainment and this value was considered the initial concentration of dye. After irradiation, the MB dye degradation was 47.21% at the first 15 min of irradiation, which increased gradually and reached 97.67% after 210 min of irradiation. The initial sharp increase in dye degradation results from dye adsorption. Over time, degradation products compete with dye ions for adsorption on the catalyst surface, slowing down the photocatalytic degradation process. The

kinetics of MB dye degradation, assessed through first-order and second-order models, are depicted in Figs. S2-S3. Photocatalytic dye degradation is significantly influenced by catalyst surface adsorption and subsequent degradation by reactive species. The MB dye degradation follows pseudo-first-order kinetics (Fig. S2). The mechanism of MB dye degradation under light irradiation using CuO NPs is shown in Eqs. 4-8. The electrons excitation from VB to CB produces an e^- and h^+ pair upon irradiation. The e^- (CB) reacts with O_2 and give a superoxide radical, while the h^+ reacts with H_2O to form $\cdot OH$ radical. The $\cdot OH$ radicals are reactive species that degrade the MO dye oxidatively into a low molecular weight intermediate and, finally, into H_2O , CO_2 , and inorganic ions (Eqs. 4-8). The proposed MB dye degradation mechanism using CuO NPs prepared via green route is depicted in Fig. 8. PCA of CuO NPs shows potential for cost-effective MO dye removal under light, offering an economical solution for MB dye remediation, which is also in line with previous studies (Table 2). Green synthesis, using natural compounds like plant extracts, microorganisms, and green reducing agents, is an eco-friendly approach with catalytic applications. Consideration of plant extract and extraction method is crucial for tailoring nanoparticles to specific properties and applications (Sinha and Ahmaruzzaman 2015, Wang et al., 2019, Kusuma et al., 2023).



3.7. Reusability and recyclability of the catalyst

The development of green technologies is a key focus in industrial discharged water treatment strategies, with a focus on the repeated use of photocatalysts because of their advantages in

terms of both cost and environmental impact. The reusability of the catalyst is assessed over the course of six cycles. The CuONP was separated from the reaction mixture of MB dye degradation, washed with distilled water, and dried before being added to a new dye solution for further cycles of photodegradation in order to evaluate its reusability. MB degradation was 97.67%¹⁷ in the first cycle, declined slightly to 96.1% in the second cycle, and showed 94.6% degradation in the third cycle. In the sixth cycle, the MB dye degradation decreased even more, reaching 89.9% (Fig. S4). The decline in catalytic activity is attributed to dye accumulation on active sites. Reusable catalysts in photocatalytic processes offer environmental and economic sustainability by reducing resource consumption, waste, and overall environmental impact. Single-use catalysts, especially those with hazardous or difficult-to-recycle components, pose disposal challenges, making reusability a more sustainable choice.

4. Conclusion

¹ The synthesis of copper oxide nanoparticles (CuO NPs) utilized *Jasminum sambac* leaf extract, wherein nanoparticle characteristics were notably influenced by solution pH, temperature, reaction time, and copper sulfate concentration. The resulting CuO NPs were monodisperse, crystalline, monoclinic, and exhibited purity, with a particle size of 13.4 nm. These NPs displayed promising photocatalytic activity, achieving a notable 97% removal of Methylene Blue (MB) dye after 210 minutes of light exposure. Factors such as pH, CuO NPs dosage, and MB dye concentration influenced the photocatalytic performance. The green synthesis method using *Jasminum sambac* leaf extract proves highly efficient for tailoring CuO NPs for photocatalytic applications. Future research will focus on evaluating photocatalytic activity under visible light and studying the toxicity of treated MB dye, including the identification of intermediate byproducts.

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