

## Bio-fabricated CuO NPs as green catalyst towards remediation of environmental pollutants

Boya Palajonnala Narasaiah<sup>1</sup>, Badal Kumar Mandal<sup>1,\*</sup> <sup>1</sup>Trace Elements Speciation Research Laboratory, Department of Chemistry, School of Advanced Sciences, Vellore Institute of Technology (VIT), Vellore 632014, Tamil Nadu, India\*corresponding author e-mail address: [badalmandal@vit.ac.in](mailto:badalmandal@vit.ac.in); [badalkmandal@gmail.com](mailto:badalkmandal@gmail.com) | Scopus ID: [7102080708](https://orcid.org/7102080708)

## ABSTRACT

We report the fabrication of biogenic CuO NPs by green chemistry concept using agriculture waste durva grass extract as stabilizing and capping agent. The green methods offer few advantageous over the common physical and chemicals procedures due to easy, fast and eco-friendly i.e. without using costly and hazardous chemicals. The synthesized CuO NPs were characterized using different instrumental techniques such as UV-Vis, IR, XRD and TEM. UV-Vis spectroscopy confirmed the presence of CuO NPs by SPR peak at 299 nm. Fourier transform infrared spectroscopy (FTIR) identified the characteristic functional groups of synthesised CuO NPs as well as extract and Powder-XRD conformed for the crystalline nature of CuO NPs. The morphology of the synthesized CuO NPs was monitored by transmission electron microscopy (TEM). In addition, we have performed photocatalytic activity towards pollutant dyes such as Rhodamine B and Congo red. Our study results suggest that the synthesized CuO NPs exhibited good catalytic activity and it could be promising material for the removal/degradation of pollutants dye from waste water.

**Keywords:** *Agro-waste Durva grass, CuO NPs, Photocatalytic activity, Pollutant Dyes; Rhodamine B, Congo red.*

## 1. INTRODUCTION

In the last decades nanoparticles (NPs) and nanocatalysts have become very promising materials in nanochemistry, because of smaller size with high surface area favors enhanced catalytic activity with respect to bulk materials. Predominantly copper oxide NPs (CuO NPs) has various applications in the nanotechnology due to their good catalytic activity and antibacterial properties [1, 2]. The properties of NPs mainly depend on their size, shape, structure, composition and morphologies [3]. These tiny particles have been synthesized by physical, chemical and biological methods [4, 5, 6]. These are expensive methods and require high temperature, pressure and energy since other reagents and solvents used are potentially hazardous to the environment and biological systems. Hence an ecofriendly process for the synthesis of NPs which does not use hazardous toxic chemicals is needed urgently [7, 8]. Synthesis of NPs using plants as a possible alternative method to physical and chemical process are considered due to environmentally eco-friendly and using nontoxic solvents and reagents [9]. The synthesis of metal NPs using various plants extracts including watermelon rind is reported [10]. The uses of CuO NPs are very important for its incredible and selective catalytic activity [11].

Nowadays residual organic dyes in effluents of different industries such as textiles, paper, paints, printing, plastic, pharmaceutical, cosmetic, medicine, food industries are a global concern. The textile industries play a major role in the economy of the USA and the associated countries. It accounts for the largest spending of dyestuffs @ 80% taking overall account, which amounts about to 80000 tons in India. India ranks second after China in the export business of dyestuff. Actually, about  $10^5$  -  $10^6$  tons of synthetic dyes produced annually are released into the environment as wastewaters globally. Being hazardous to the environment even 1 mg/L of dye can cause serious problems to environments. Dyes

contaminated wastewater is toxic, poisonous, dangerous, and carcinogenic to living beings [12]. Hence the international environmental standards have become more and more technological dependent for the removal of organic pollutants such as dyes. Thus the remediation of all unwanted dyes i.e. Methylene blue (MB), Congo red (CR), Rhodamine B (Rh-B) and Methyl orange (MO) in waste water is a challenging task to the environmentalists.

The various methods accessible for the degradation of organic pollutants such as biological, physical, and chemical techniques including aerobic, anaerobic, coagulation, chemical oxidation, membrane separation, electrochemical treatment, dilution, filtration, ion exchange and RO membranes are active in waste water treatments with regard to removal of dye stuff [13-15]. Many techniques have been widely applied in wastewater treatments but they are associated with some drawbacks for their applications. Hence safe and cheap technologies are highly preferable for remediation. Recently enhanced catalytic degradation of pollutants finds some importance in the treatment of wastewater generated from industrial and living sources [16, 17]. Among different catalysts nanocatalysts have undergone enormous growth in recent years and seem to be an emerging material in the field of catalysis in the near future [18].

The present study focused on the biosynthesis of copper oxide nanoparticles (CuO NPs) from agro-waste durva grass aqueous extract for the first-time. The green-synthesized CuO NPs was used for the degradation of environmentally harmful dyes. Especially the ability of the synthesized CuO NPs as an efficient green photocatalyst in degrading Rhodamine B and Congo red was investigated by UV irradiation at room temperature in the present study.

## 2. EXPERIMENTAL SECTION

### Materials.

Cu (NO<sub>3</sub>)<sub>2</sub> · 3H<sub>2</sub>O, Rhodamine B (RB), and Congo red (CR) were purchased from Sigma-Aldrich, India. Distilled water was used as a solvent and all analytical grade reagents were used. Fresh durva grass was collected from the agriculture crop field of AP state, India.

### Collection and Preparation of Durva Grass Aqueous Extract.

The green durva grass was collected from sunflower crop cultivated agricultural land. After washing thoroughly durva grass was cut into small pieces and washed with hot water repeatedly to remove soluble matters, then dried in an oven at 60 °C for 3h. The oven-dried powder was sieved through 100 mesh sieve. 100 mL of distilled water containing 3 g powder was heated in a water-bath at 80 °C for 30 min to prepare aqueous extract. The extract was cooled, filtered through Whatman filter paper and was stored for the synthesis of CuO NPs.

### Synthesis of Copper Oxide Nanoparticles.

0.3624 g copper nitrate (0.05 M) was dissolved in 30 mL double distilled water and then 30 mL of freshly prepared aqueous extract was added dropwise with stirring thoroughly at 70 °C for 3h. The colour change from light green to dark brown during the heating process was due to the surface Plasmon resonance of CuO NPs formed which was further confirmed by UV-Visible spectroscopy. The obtained colloidal solution was centrifuged at 5000 rpm for 30 min, then black precipitates were washed thrice with absolute ethanol followed by double distilled water to remove any biomolecules and impurities present. The obtained fine pellet was dried on a hot plate at 60 °C for 20 min, crushed into powder and then was annealed at 500 °C 3h to get crystalline CuO NPs.

### Characterization of Synthesized CuO NPs.

The preliminary characterization was carried out by a UV-Visible spectroscopy (Jasco V-670 UV-Visible double beam spectrophotometer). The absorption intensity of the synthesized colloidal CuO NPs solution was recorded in between wavelength range of 200 to 800 nm, and the obtained data was re-plotted by using Origin 8.5 software.

FTIR analysis was carried out to detect the presence of functional groups in durva grass extract and on the surface of the synthesized CuO NPs. Separated dried CuO NPs was used for XRD and FTIR analysis. The purified dried CuO NPs was analysed using in KBr

pellets by Shimadzu IR AFFINITY-1 (JASCO FT-IR 4100) against durva grass powder as control in the diffuse reflectance mode at a resolution of 4cm<sup>-1</sup>. The XRD analysis of purified CuO NP was carried out by using a Bruker D8 Advance diffractometer with Cu K $\alpha$  radiation ( $\lambda=1.54$  Å, scanning range (2 $\theta$  value) of 10° to 90°, scanning rate of 4° /min and step size of 0.02°).

The high resolution transmission electron microscopy (HR-TEM) checked the size and shape of the synthesized CuO NPs. The selected area electron diffraction (SAED) pattern was recorded using a JEOL-JEM 2100 transmission electron microscope with an acceleration voltage of 200 kV. Initially 15 mg CuO NPs pure solid powder was dispersed in 2 mL absolute ethanol by sonication for 10 min using an ultrasonic bath for microscopic analysis. The TEM samples of CuO NPs were prepared by placing a drop of solution on the lacey carbon coated copper grids followed by evaporation of solvents from grids at ambient conditions for TEM analysis.

Dynamic light scattering (DLS) study was carried out to determine Zeta potential in predicting the stability of CuO NPs in aqueous dispersion by using Horiba scientific nanoparticci (nanoparticles analyzer, SZ-100). The suspension was measured in cell culture medium at 25 °C and 150 V in triplet for each sample. Also, polydispersity indexes (PDI) was determined at 20 °C with a scattering angle of 173° in triplicate. Deionized water was used as a source of dispersing medium, while standard Malvern-50 v was used to calibrate instrument before analysis of samples.

### Photocatalytic Activity of CuO NPs.

The photocatalytic activity of synthesized CuO NPs was carried out in an aqueous medium against degradation of organic dyes Rhodamine B (Rh-B) and Congo red (CR) by using Heber multilamp photoreactor under UV irradiation ( $\lambda_{max}$  254 nm) at 125 W and low-pressure. 60 mL dye (10 mg/L) in 100 mL capacity of quartz tube was continuously stirred under dark in the absence of catalyst; the variation in absorption intensity was monitored by UV-Visible spectroscopy. The second phototube containing 60 mL dye solution (10 mg/L) with different doses of CuO NPs (10, 20, 30 mg) was irradiated with UV light inside photoreactor. The photodegradation of organic dyes was observed by measuring absorption intensity of the dye solution at regular time intervals within  $\lambda_{max}$  of 200-800 nm.

## 3. RESULTS SECTION

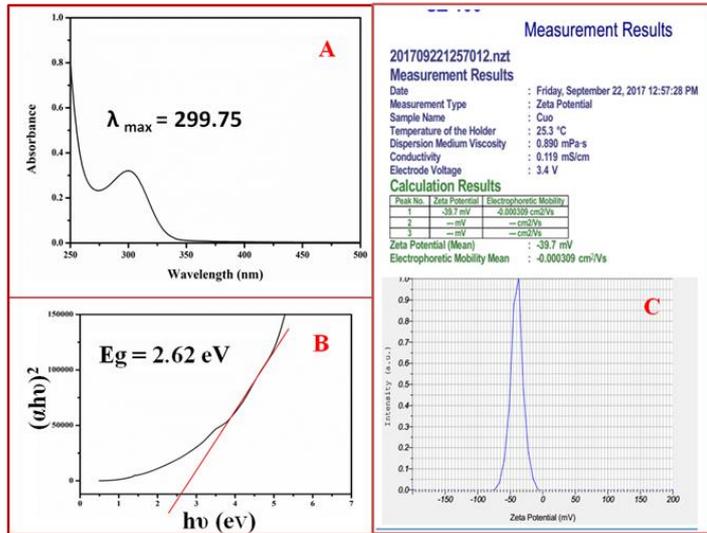
Initially the formation of CuO NPs indicated colour change from light yellow to dark brown and also the formation of CuO NPs was confirmed by UV-Visible spectroscopy where the absorbance of the NPs dispersion was monitored at 200 nm to 800 nm. Fig. 1(A) shows the clear surface Plasmon resonance (SPR) band at 299.75 nm which confirms the formation of CuO NPs. The photocatalytic activity of CuO NPs depends upon its optical band gap energy. Optical band gap energy of the synthesized CuO NPs was determined by Tauc equation (i).

$$\alpha = C (h\nu - E_g)^2 \dots\dots(i)$$

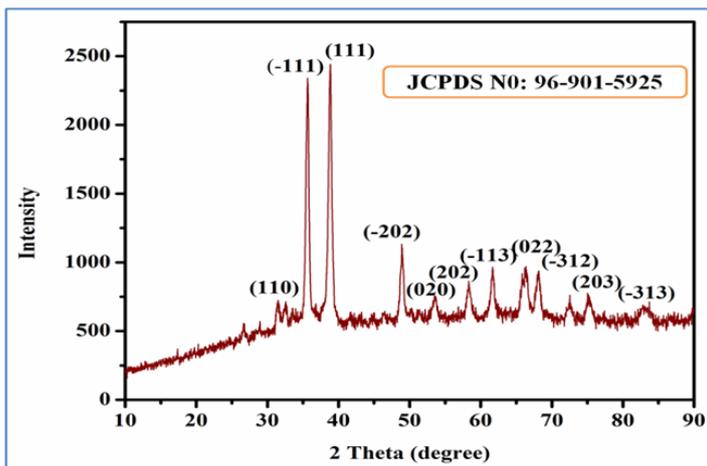
where,  $\alpha$  is the absorption coefficient, C is the constant,  $h\nu$  is the photon energy and  $E_g$  is the optical band gap energy respectively. Fig. 1(B) shows the UV-Vis DRS plot of the synthesized CuO NPs where a plot is drawn keeping  $(\alpha h\nu)^2$  as Y-axis vs.  $h\nu$  as X-axis. The band gap energy ( $E_g$ ) of CuO NPs is calculated by using the formula (i) as 2.62 eV which suggests its semiconducting property.

The measurement of zeta potential value was determined by using photon correlation spectroscopy (PCS) i.e. dynamic light scattering (DLS) study. The zeta potential value of CuO NPs'

dispersion was -39.7 mV. The high negative value indicates that CuO NPs repelled to each other in dispersion resulting high electrophoretic mobility of NPs which prevented aggregation/agglomeration of NPs [19]. The higher zeta potential values result in higher dispersion or more stability in solution (Fig. 1C).



**Figure 1.** UV-Visible absorption spectrum of the synthesized CuO NPs (A), UV-DRS plot of CuO NPs (B), DLS spectrum of CuO NPs for Zeta potential analysis (C).



**Figure 2.** XRD pattern of the synthesized CuO NPs using durva grass aqueous extract.

X-ray diffraction study was performed to get information on crystallinity and phase formation as well as purity of nanomaterials. X-ray diffraction pattern of the synthesized CuO NPs is shown in Fig. 2. XRD pattern shows characteristic reflections peaks at hkl values of (110), (-111), ((111), (-202), (020), (202), (-113), (022), (203), (-313) suggesting FCC crystal structure with lattice parameter “a” of 4.09 Å [20]. The (111) lattice plane is more intense because of its predominant orientation than other peaks and the broaden Bragg’s peaks indicate the formation of smaller crystallites. The obtained XRD pattern is compared with JCPDS data card (JCPDS N0: 96-901-5925) which confirms crystalline nature of the synthesized CuO NPs.

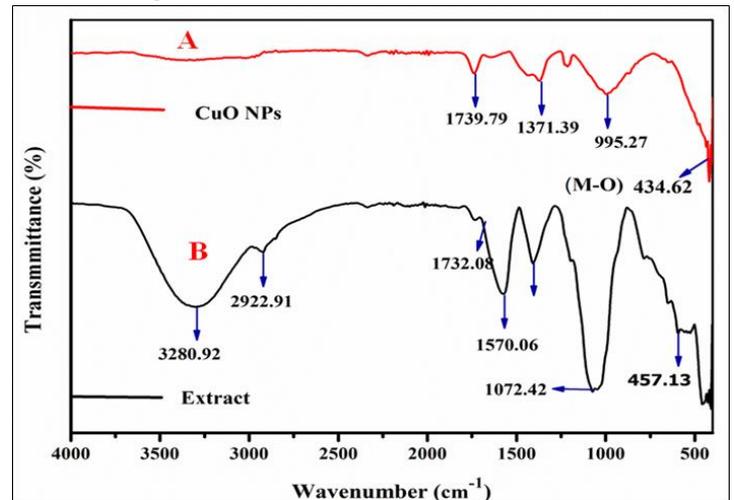
The average crystallite size is calculated by using Scherrer equation as

$$D = K\lambda/\beta\cos(\theta) \dots\dots(ii)$$

Where D is the average crystal size,  $\lambda$  is X-ray wave length ( $\lambda = 1.5406 \text{ \AA}$ ), K is the Scherrer coefficient (0.891),  $\beta$  is the full

width at half maximum intensity (FWHM) in radians, and  $\theta$  is Bragg’s angle ( $2\theta$ ). The average crystallite size was 26 nm.

Fig. 3(A) shows the FTIR spectrum of the synthesized CuO NPs and durva grass aqueous extract was also analysed to investigate biomolecules present in durva grass extract responsible for the synthesis of CuO NPs and stabilization of NPs via surface capping of biomolecules on its surfaces [21]. FTIR spectrum (Fig. 3B) of the plant extracts shows predominant bands at 3280.92, 2922.91, 1732.08, 1570.06 cm<sup>-1</sup>, 1072.42, and 457.13 cm<sup>-1</sup>. The broad band around 3100-3600 cm<sup>-1</sup> corresponds to -OH stretching frequency of polyphenolic compounds, glucose or NH- stretching of amino acid and proteins. The band at 2850-2950 cm<sup>-1</sup> corresponds to alkyl C-H stretching frequency, band at 1732.08 cm<sup>-1</sup> corresponds to carbonyl of carbohydrates. The band at 434.62 cm<sup>-1</sup> corresponds to metal-oxygen (Cu-O) stretching vibration of CuO NPs (Fig. 3A).



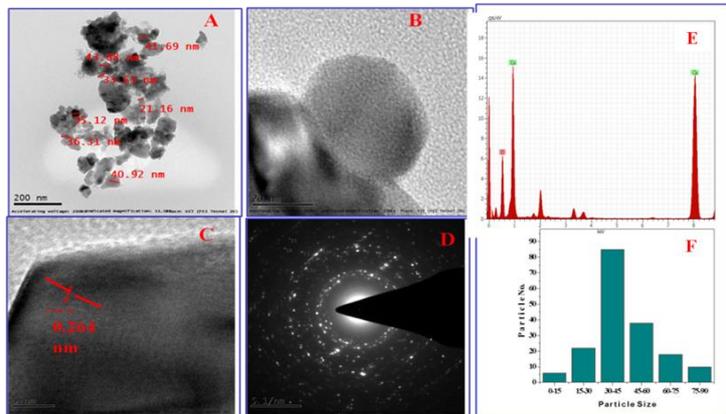
**Figure 3.** FTIR spectra of the synthesized CuO NPs (A) and durva grass aqueous extract (B).

Fig. 4 demonstrates the TEM, HR-TEM, SAED pattern and EDX (energy dispersive X-ray) micrographs of the synthesized CuO NPs. Fig. 4(A) exhibits particles size distribution and the majority of CuO NPs was within 30-45 nm with a mean diameter of 35 nm which supports the crystallite/grain size obtained from the XRD diffraction pattern by using Scherrer’s equation (Fig. 4B). Fig. 4(C) shows the inter-planar distance i.e. d-spacing value of 0.264 nm which corresponds to lattice spacing (111) plane. Fig. 4(D) shows SAED pattern where bright spots are arranged in the form of concentric circles indicating the polycrystalline nature of CuO NPs [22]. Fig. 4(E) shows EDX spectrum of CuO NPs which basically check the purity of the NPs by analyzing chemical composition of the sample. It clearly shows that the atomic % and weight % of Copper are 56.01 and 43.02; atomic % and weight % of Oxygen are 83.49 and 16.51 respectively. The above results suggest that CuO NPs are pure and polycrystalline in nature. Fig. 4(F) shows the particles size distribution within the range of 15-90 nm with a mean diameter of 35 nm.

### Photocatalytic Degradation of Dyes.

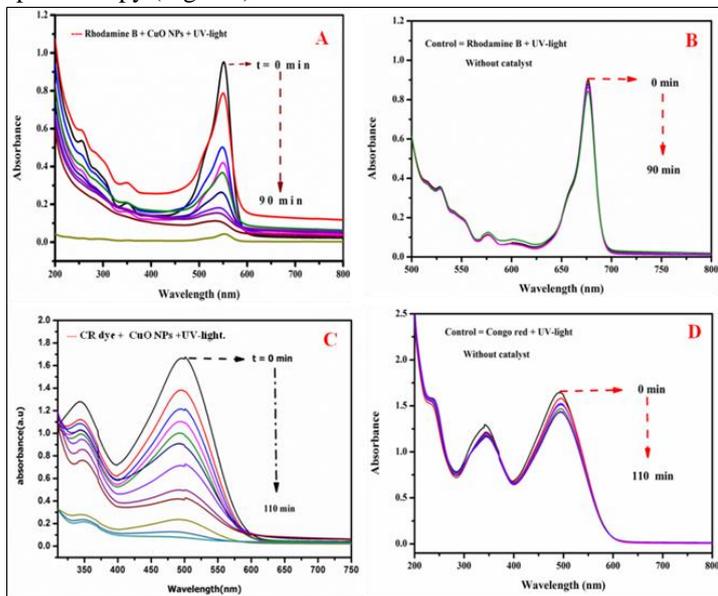
Several industries discharge a huge amount of toxic chemicals including coloring agents which are light sensitive. Among all colouring agents, Rh-B and CR are more stable organic substances with heterocyclic aromatic rings [23, 24]. Heterocyclic

aromatic rings exhibit strong resistance to heat and light. Environmentally nanoparticles are considered to be less toxic and have high photocatalytic activity with large surface areas. Among different M/MO NPs (Table 1), CuO NPs with high photocatalytic activity is studied extensively towards degradation of Rh-B and CR dyes under UV irradiation.



**Figure 4.** HR-TEM micrographs of CuO NPs at 200 nm magnification (A), 20 nm magnification (B), 5 nm magnification (C), SAED pattern of CuO NPs (D), EDX spectrum of CuO NPs (E) and particles size distribution of CuO NPs by DLS (F).

The photocatalytic degradation of Rh-B and CR dyes was evaluated by the photocatalyst CuO NPs in a photoreactor fitted with UV multilamp (125 W,  $\lambda_{max}$  254 nm). 20 mg CuO NPs was added to 60 mL dye (10 mg/L) solution with stirring for 30 min at room temperature to reach adsorption-desorption equilibrium under UV irradiation (125 W, 254 nm). All dye samples were collected every 10 min intervals after UV irradiation for 90 min until colorless solution of Rh-B dye and monitored by UV-Vis spectroscopy (Fig. 5A).

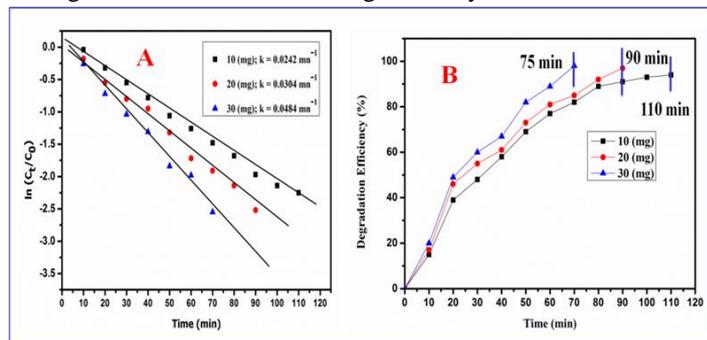


**Figure 5.** Degradation of Rh-B dye by catalyst CuO NPs (A) and without catalyst CuO NPs (B), degradation of CR dye by catalyst CuO NPs (C) and without catalyst (D) under UV irradiation.

The dye degradation was monitored by UV-Visible spectroscopy at 665 nm for Rh-B and at 556 nm for CR dye respectively. All samples were collected every 10 min intervals up to 110 min until a complete colorless solution of CR dye (Fig. 5C). Also, the degradation of dyes was measured by UV-Visible spectroscopy

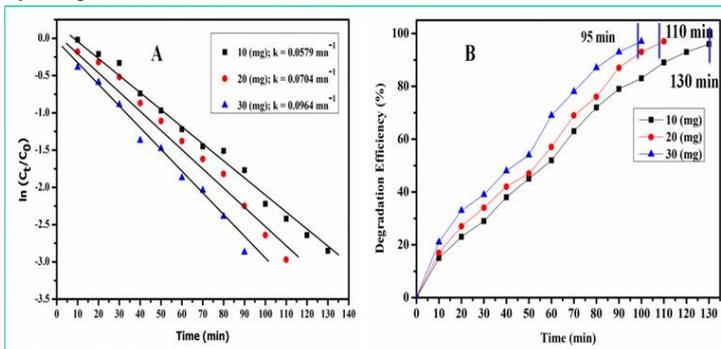
without catalyst as the control sample. There was no change in absorbance value of dye without catalyst CuO NPs (Fig. 5B&D).

Fig. 6(A&B) shows the effect of catalyst doses (10, 20, and 30 mg) on degradation kinetics of dyes. The degradation rate of Rh-B and CR dyes was increased with an increase in catalyst dose [25, 26]. Also, degradation efficiency was increased i.e. time taken for degradation decreased with higher catalyst dose.



**Figure 6.** Effect of catalyst dose on degradation of Rh-B dye: First order linear plot of  $\ln(C_t/C_0)$  vs. time (min) (A), degradation efficiency (%) of Rh-B dye (B).

The degradation efficiency of Rh-B and CR was calculated by using the following formula as Degradation of Dye (%) =  $[(C_0 - C_t) * 100 / C_0]$ , where  $C_0$  and  $C_t$  are a concentration of dye at time 'zero' and time 't' respectively. The obtained results show that 98% of Rh-B degraded within 90 min and 95% of CR degraded within 110 min which suggest that CR is a stronger organic dye than Rh-B i.e. the catalyst CuO NPs degraded Rh-B quickly compare to CR degradation at a particular dose of catalyst. The obtained results are compared with other reported literatures on dye degradation (Table 1).



**Figure 7.** Effect of catalyst dose on photo-degradation of CR dye: First order linear plot of  $\ln(C_t/C_0)$  vs. time (A), degradation efficiency (%) of CR dye (B).

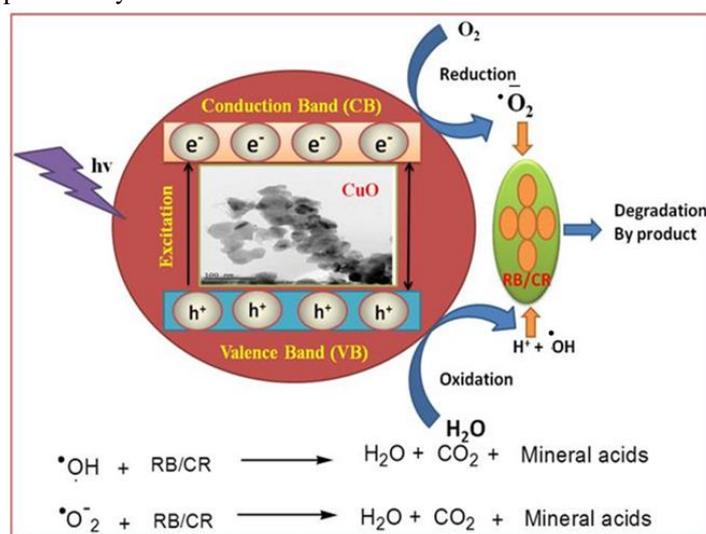
The degradation efficiency of NPs depends on its size, surface area and morphology [27]. The degradation rate of dye was increased with NPs having lower band gap value which can readily generate electron-hole pairs easily available for photodegradation on UV irradiation like CuO NPs. Hole can highly oxidise the organic molecules into carbon dioxide and water molecules; while the excited electrons are scavenging by 'O<sub>2</sub>' in the environment forming super oxide which is a very strong oxidising agent. The photocatalytic degradation of dyes by photocatalyst CuO NPs depends on three important parameters i.e. size of CuO NPs, bandgap energy and nature of dyes. Normally, more number of aromatic ring, hetero-aromatic rings and azo dyes are strong/stable dyes such as CR dye which is stronger dye than Rh-B dye [28].

The photocatalytic activity CuO NPs follows the first order reaction in degrading CR and Rh-B dyes and the rate of degradation is calculated by the following formula [29] as  $\ln(C_t) = \ln(C_0) - kt$ , where,  $C_0$  is the initial concentration of dye,  $C_t$  is the final concentration of dye at time 't' and k is the rate constant, respectively. Fig. 7 shows a graphical plot between  $\ln C_t/C_0$  along Y-axis vs. time (min) along X-axis suggesting a decreasing trend under UV-irradiation for degradation of dyes. The rate constant 'k' of Rh-B and CR photodegradation was checked from the plots which are 0.0579, 0.0704 and 0.0964  $\text{min}^{-1}$  for 10, 20 and 30 mg of catalyst mediated photodegradation, respectively (Fig. 7 A&B). The obtained results clearly show that the value of rate constant was increased with increasing catalyst doses (Table 2).

#### Photocatalytic Degradation Mechanism of Dye.

Fig. 8 shows a plausible mechanism on photocatalytic degradation of Rh-B/CR dye using CuO NPs as photocatalyst under UV-irradiation. In the photodegradation process CuO NPs mediates reduction of oxygen to generate super oxide ( $\text{O}_2^-$ ) using excited conduction band electrons under UV-irradiation, while positive holes on valence band help to generate another reactive oxygen species hydroxyl radical ' $\text{OH}^\bullet$ ' after reduction of water

molecules which causes photodegradation of toxic dyes [30]. The present results suggest degradation of toxic dyes to water molecules and carbon dioxide in the presence of CuO NPs as photocatalyst.



**Figure 8.** Possible mechanism on photocatalytic degradation of RB/CR dye using CuO NPs as photocatalyst.

## 4. CONCLUSIONS

Our study reported the biosynthesis of CuO NPs by argo-waste durva grass aqueous extract for the first time in literature. The synthesised CuO NPs was confirmed by UV-Visible, FTIR, XRD, HR-TEM and EDX while DLS was used for the stability study. Under UV-irradiation the synthesized CuO NPs showed

good photocatalytic efficiency towards the degradation of CR and Rh-B dyes. The current study reports that the green synthesized CuO NPs could be used as an effective photocatalyst in degrading environmentally hazardous and toxic dyes in a greener way.

**Table 1.** Comparison of degradation efficiency of synthesized CuO NPs with other published reports on dye degradation.

Dye name	Photo-catalyst	Time (min)	Deg. (%)	References
<b>Rh-B</b>	CuO NPs	300	100	[31]
	CuO-nanowires	660	95	[32]
	CuO/ZnO NPs	150	83	[33]
	CuO nanostructures	300	85	[34]
	Fe <sub>3</sub> O <sub>4</sub> /Zn-/CuWO <sub>4</sub> NPs	210	99	[35]
	CuO NPs	90	98	Present study
<b>CR</b>	Ba/Alg/CMC/TiO <sub>2</sub>	240	91	[36]
	ZnO NPs	135	97	[37]
	Chitosan/nano-CdS	180	85.9	[38]
	WO <sub>3</sub> /TiO <sub>2</sub> /AC	120	95.21	[39]
	CeO <sub>2</sub> NPs	100	95	[40]
	CuO NPs	110	95	Present study

**Table 2.** Effect of CuO NPs as photocatalyst on degradation of Rh-B and CR dyes.

Dye	Catalyst (mg)	Con. Dye (mg/L)	Degradation Time (min)	% of degradation	Rate Constant k ( $\text{min}^{-1}$ )
<b>Rh-B</b>	10	10	110	96	0.0242
	20	10	90	98	0.0304
	30	10	75	98	0.0484
<b>CR</b>	10	10	130	93	0.0579
	20	10	110	95	0.0704
	30	10	95	96	0.0964

## 5. REFERENCES

- Jin, Z.; Liu, C.; Qi, K.; Cui, X. Photo-reduced Cu/CuO nanoclusters on TiO<sub>2</sub> nanotube arrays as highly efficient and reusable catalyst. *Sci. Rep.* **2017**, *7*, 39695, <https://doi.org/10.1038/srep39695>.
- Azam, A.; Ahmed, A.S.; Oves, M.; Khan, M.S.; Memic, A. Size-dependent antimicrobial properties of CuO nanoparticles against Gram-positive and-negative bacterial strains. *Int. J. Nanomed.* **2012**, *7*, 3527, <https://doi.org/10.2147/IJN.S29020>.

3. Cuenya, B.R. Synthesis and catalytic properties of metal nanoparticles: Size, shape, support, composition, and oxidation state effects. *Thin Solid Films* **2010**, *518*, 3127-3150, <https://doi.org/10.1016/j.tsf.2010.01.018>.
4. Padovini, D.S.S.; Pontes, D.S.L.; Dalmascio, C.J.; Pontes, F.M.; Longo, E. Facile synthesis and characterization of ZrO<sub>2</sub> nanoparticles prepared by the AOP/hydrothermal route. *Rsc Adv.* **2014**, *4*, 38484-38490, <https://doi.org/10.1039/C4RA04861J>.
5. Coker, V.S.; Bennett, J.A.; Telling, N.D.; Henkel, T.; Charnock, J.M.; Van der Laan, G.; Patrick, R.A.; Pearce, C.I.; Cutting, R.S.; Shannon, I.J.; Wood, J. Microbial engineering of nanoheterostructures: biological synthesis of a magnetically recoverable palladium nanocatalyst. *ACS Nano* **2010**, *4*, 2577-2584, <https://doi.org/10.1021/nm9017944>.
6. Li, M.; He, W.; Liu, Y.; Wu, H.; Wamer, W.G.; Lo, Y.M.; Yin, J.J.; FD&C Yellow No. 5 (Tartrazine) degradation via reactive oxygen species triggered by TiO<sub>2</sub> and Au/TiO<sub>2</sub> nanoparticles exposed to simulated sunlight. *J. Agric. Food Chem.* **2014**, *62*, 12052-12060, <https://doi.org/10.1021/jf5045052>.
7. Habeeb, M.K. Biosynthesis of nanoparticles by microorganisms and their applications. *Int. J. Adv. Sci. Technol. Res.* **2013**, *1*, 44-51, <http://dx.doi.org/10.1155/2011/270974>.
8. Akhtar, M.S.; Panwar, J.; Yun, Y.S. Biogenic synthesis of metallic nanoparticles by plant extracts. *ACS Sustain Chem. Eng.* **2013**, *1*, 591-602, <https://doi.org/10.1021/sc300118u>.
9. Shamaila, S.; Sajjad, A.K.L.; Farooqi, S.A.; Jabeen, N.; Majeed, S.; Farooq, I. Advancements in nanoparticle fabrication by hazard free eco-friendly green routes. *Appl. Mater. Today.* **2016**, *5*, 150-199, <https://doi.org/10.1016/j.apmt.2016.09.009>.
10. Lakshmipathy, R.; Reddy, B.P.; Sarada, N.C.; Chidambaram, K.; Pasha, S.K. Watermelon rind-mediated green synthesis of noble palladium nanoparticles: catalytic application. *Appl. Nanosci.* **2015**, *5*, 223-228, <https://doi.org/10.1007/s13204-014-0309-2>.
11. Jin, Z.; Liu, C.; Qi, K.; Cui, X. Photo-reduced Cu/CuO nanoclusters on TiO<sub>2</sub> nanotube arrays as highly efficient and reusable catalyst. *Sci. Rep.* **2017**, *7*, 39695, <https://doi.org/10.1038/srep39695>.
12. Kant, R. Textile dyeing industry an environmental hazard. *Natural Sci.* **2012**, *4*, 22-26, <https://doi.org/10.4236/ns.2012.41004>.
13. Khodadadi, B.; Bordbar, M.; Nasrollahzadeh, M. Green synthesis of Pd nanoparticles at Apricot kernel shell substrate using Salvia hydrangea extract: catalytic activity for reduction of organic dyes. *J. Colloid Interface Sci.* **2017**, *490*, 1-10, <https://doi.org/10.1016/j.jcis.2016.11.032>.
14. Bhattacharjee, A.; Ahmaruzzaman, M. A novel and green process for the production of tin oxide quantum dots and its application as a photocatalyst for the degradation of dyes from aqueous phase. *J. Colloid Interface Sci.* **2015**, *448*, 130-139, <https://doi.org/10.1016/j.jcis.2015.01.083>.
15. Ajmal, A.; Majeed, I.; Malik, R.N.; Idriss, H.; Nadeem, M.A. Principles and mechanisms of photocatalytic dye degradation on TiO<sub>2</sub> based photocatalysts: a comparative overview. *Rsc Adv.* **2014**, *4*, 37003-37026, <https://doi.org/10.1039/C4RA06658H>.
16. Sreelatha, S.; Velvizhi, G.; Reddy, C.N.; Modestra, J.A.; Mohan, S.V. Solid electron acceptor effect on biocatalyst activity in treating azo dye based wastewater. *RSC Adv.* **2015**, *5*, 95926-95938, <https://doi.org/10.1039/c5ra15648c>.
17. Ahmad, A.; Mohd-Setapar, S.H.; Chuong, C.S.; Khatoun, A.; Wani, W.A.; Kumar, R.; Rafatullah, M. Recent advances in new generation dye removal technologies: novel search for approaches to reprocess wastewater. *RSC Adv.* **2015**, *5*, 30801-30818, <https://doi.org/10.1039/C4RA16959J>.
18. Gao, J.F.; Li, H.Y.; Pan, K.L.; Si, C.Y. Green synthesis of nanoscale zero-valent iron using a grape seed extract as a stabilizing agent and the application for quick decolorization of azo and anthraquinone dyes. *RSC Adv.* **2016**, *6*, 22526-22537, <https://doi.org/10.1039/C5RA26668H>.
19. Kumar, K.M.; Mandal, B.K.; Tammina, S.K. Green synthesis of nano platinum using naturally occurring polyphenols. *RSC Adv.* **2013**, *3*, 4033-4039, <https://doi.org/10.1039/C3RA22959A>.
20. Sadollahkhani, A.; Ibupoto, Z.H.; Elhag, S.; Nur, O.; Willander, M. Photocatalytic properties of different morphologies of CuO for the degradation of Congo red organic dye. *Ceram. Int.* **2014**, *40*, 11311-11317, <https://doi.org/10.1016/j.ceramint.2014.03.132>.
21. Tamuly, C.; Saikia, I.; Hazarika, M.; Das, M.R. Reduction of aromatic nitro compounds catalyzed by biogenic CuO nanoparticles. *RSC Adv.* **2014**, *4*, 53229-53236, <https://doi.org/10.1039/C4RA10397A>.
22. Mott, D.; Galkowski, J.; Wang, L.; Luo, J.; Zhong, C.J. Synthesis of size-controlled and shaped copper nanoparticles. *Langmuir* **2007**, *23*, 5740-5745, <https://doi.org/10.1021/la0635092>.
23. Malwal, D.; Gopinath, P. Enhanced photocatalytic activity of hierarchical three dimensional metal oxide@ CuO nanostructures towards the degradation of Congo red dye under solar radiation. *Catal. Sci. Technol.* **2016**, *6*, 4458-4472, <https://doi.org/10.1039/C6CY00128A>.
24. Rahman, Q.I.; Ahmad, M.; Misra, S.K.; Lohani, M. Effective photocatalytic degradation of rhodamine B dye by ZnO nanoparticles. *Mater. Lett.* **2013**, *91*, 170-174, <https://doi.org/10.1016/j.matlet.2012.09.044>.
25. Tammina, S.K.; Mandal, B.K. Tyrosine mediated synthesis of SnO<sub>2</sub> nanoparticles and their photocatalytic activity towards Violet 4 BSN dye. *J. Mol. Liq.* **2016**, *221*, 415-421, <https://doi.org/10.1016/j.molliq.2016.05.079>.
26. Gautam, A.; Kshirsagar, A.; Biswas, R.; Banerjee, S.; Khanna, P.K.; Photodegradation of organic dyes based on anatase and rutile TiO<sub>2</sub> nanoparticles. *RSC Adv.* **2016**, *6*, 2746-2759, <https://doi.org/10.1039/C5RA20861K>.
27. Mao, B.; An, Q.; Zhai, B.; Xiao, Z.; Zhai, S. Multifunctional hollow polydopamine-based composites (Fe<sub>3</sub>O<sub>4</sub>/PDA@Ag) for efficient degradation of organic dyes. *RSC Adv.* **2016**, *6*, 47761-47770, <https://doi.org/10.1039/C6RA05954F>.
28. Kirankumar, V.S.; Sumathi, S. Structural, optical, magnetic and photocatalytic properties of bismuth doped copper aluminate nanoparticles. *Mater. Chem. Phys.* **2017**, *197*, 17-26, <https://doi.org/10.1016/j.matchemphys.2017.05.021>.
29. Khan, S.S. Enhancement of visible light photocatalytic activity of CdO modified ZnO nanohybrid particles. *J. Photochem. Photobiol.* **2015**, *142*, 1-7, <https://doi.org/10.1016/j.jphotobiol.2014.11.001>.
30. Naraginti, S.; Thejaswini, T.V.L.; Prabhakaran, D.; Sivakumar, A.; Satyanarayana, V.S.V.; Prasad, A.A. Enhanced photo-catalytic activity of Sr and Ag co-doped TiO<sub>2</sub> nanoparticles for the degradation of Direct Green-6 and Reactive Blue-160 under UV & visible light. *Spectrochim. Act. A.* **2015**, *149*, 571-579, <https://doi.org/10.1016/j.saa.2015.04.101>.
31. Tadjarodi, A.; Akhavan, O.; Bijanzad, K. Photocatalytic activity of CuO nanoparticles incorporated in mesoporous structure prepared from bis (2-aminonicotinato) copper (II) microflakes. *Trans. Nonferrous Met. Soc. China.* **2015**, *25*, 3634-3642, [https://doi.org/10.1016/S1003-6326\(15\)64004-3](https://doi.org/10.1016/S1003-6326(15)64004-3).
32. Wang, S.L.; Li, P.G.; Zhu, H.W.; Tang, W.H. Controllable synthesis and photocatalytic property of uniform CuO/Cu<sub>2</sub>O composite hollow microspheres. *Adv. Powder Technol.* **2012**, *230*, 48-53, <https://doi.org/10.1016/j.powtec.2012.06.051>.

33. Mehr, E.S.; Sorbiun, M.; Ramazani, A.; Fardood, S.T. Plant-mediated synthesis of zinc oxide and copper oxide nanoparticles by using *ferulago angulata* (schlecht) boiss extract and comparison of their photocatalytic degradation of Rhodamine B (RhB) under visible light irradiation. *J. Mater. Sci. Mater. Electron.* **2018**, *29*, 1333-1340, <https://doi.org/10.1007/s10854-017-8039-3>.
34. Zaman, S.; Zainelabdin, A.; Amin, G.; Nur, O.; Willander, M. Efficient catalytic effect of CuO nanostructures on the degradation of organic dyes. *J. Phys. Chem. Solids.* **2012**, *73*, 1320-1325, <https://doi.org/10.1016/j.jpcs.2012.07.005>.
35. Shekofteh-Gohari, M.; Habibi-Yangjeh, A. Fabrication of novel magnetically separable visible-light-driven photocatalysts through photosensitization of Fe<sub>3</sub>O<sub>4</sub>/ZnO with CuWO<sub>4</sub>. *J. Ind. Eng. Chem.* **2016**, *44*, 174-184, <https://doi.org/10.1016/j.jiec.2016.08.028>.
36. Thomas, M.; Naikoo, G.A.; Sheikh, M.U.D.; Bano, M.; Khan, F. Effective photocatalytic degradation of Congo red dye using alginate/carboxymethyl cellulose/TiO<sub>2</sub> nanocomposite hydrogel under direct sunlight irradiation. *J. Photochem. Photobiol.* **2016**, *327*, 33-43, <https://doi.org/10.1016/j.jphotochem.2016.05.005>.
37. Fowsiya, J.; Madhumitha, G.; Al-Dhabi, N.A.; Arasu, M.V. Photocatalytic degradation of Congo red using *Carissa edulis* extract capped zinc oxide nanoparticles. *J. Photochem. Photobiol.* **2016**, *162*, 395-401, <https://doi.org/10.1016/j.jphotobiol.2016.07.011>.
38. Zhu, H.; Jiang, R.; Xiao, L.; Chang, Y.; Guan, Y.; Li, X.; Zeng, G. Photocatalytic decolorization and degradation of Congo Red on innovative crosslinked chitosan/nano-CdS composite catalyst under visible light irradiation. *J. Hazard. Mater.* **2009**, *169*, 933-940, <https://doi.org/10.1016/j.jhazmat.2009.04.037>.
39. Sun, J.H.; Wang, Y.K.; Sun, R.X.; Dong, S.Y. Photodegradation of azo dye Congo Red from aqueous solution by the WO<sub>3</sub>-TiO<sub>2</sub>/activated carbon (AC) photocatalyst under the UV irradiation. *Mater. Chem. Phys.* **2009**, *115*, 303-308, <https://doi.org/10.1016/j.matchemphys.2008.12.008>.
40. Latha, P.; Prakash, K.; Karuthapandian, S. Effective Photodegradation of CR & MO dyes by morphologically controlled Cerium oxide nanocubes under visible light illumination. *Optik* **2018**, *154*, 242-250, <https://doi.org/10.1016/j.ijleo.2017.10.054>.

## 6. ACKNOWLEDGEMENTS

The authors thank Vellore Institute of Technology, Vellore for the financial support, working platform and instrument facility given to complete the study. Especially, the authors thank Vellore Institute of Technology, Vellore for providing "VIT SEED GRANT" for carrying out this research work.



© 2019 by the authors. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<http://creativecommons.org/licenses/by/4.0/>).