



Biosynthesis of copper nanoparticles using *Bacillus flexus* and estimation of their potential for decolorization of azo dyes and textile wastewater treatment

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ABSTRACT

In many underdeveloped countries, textile industries discharge their effluents without any treatment. The untreated textile wastewater consists of both azo dyes and heavy metals which not only change the physicochemical and biological properties of soil and water but also affect human health. In recent years, copper-based nano-materials have attained worldwide attention because of their unique properties and potential for decolorization of azo dyes and wastewater treatment. The present study demonstrates the bacterial synthesis of copper nanoparticles (Cu-NPs) using *Bacillus flexus* strain isolated from a textile wastewater followed by their application for photocatalytic degradation of various azo dyes and treatment of actual wastewater. The FT-IR analysis confirmed the presence of various functional groups including proteins on Cu-NPs which improved their stability. The scanning electron microscopy (SEM) images revealed the spherical shape of Cu-NPs with a size of range between 17–34 nm. Similarly, the XRD analysis of biosynthesized Cu-NPs showed various diffraction peaks at 44.5°, 51.5°, 74.75° which confirmed the crystalline nature of nanoparticles. While studying the photocatalytic decolorization of azo dyes by Cu-NPs, it was observed that 88.164 ± 0.19 %, 88.452 ± 1.89 %, 90.433 ± 1.81 %, 64.770 ± 1.02 %, 46.774 ± 1.61 %, and 67.274 ± 2.89 % of reactive black-5, congo red, malachite green, methylene blue, reactive red-2 and blue direct, respectively, were decolorized after 4 h of solar irradiation at 50 ppm concentration. Additionally, the biosynthesized nanoparticles also resulted in reduction of various parameters like EC, pH, TDS, COD, color intensity, sulphates, and phosphates in the textiles wastewater. The reduction of COD, sulfates, and phosphates was about 39.659 %, 43.157 %, and 49.493 %. The results of current work suggest that biosynthesized copper nanoparticles might serve as a potential green solution for the decolorization of various dyes including wastewater treatment.

1. Introduction

Water is one of the most important natural resource on earth not only for human beings but also for all other living organisms. All essential phenomena of the biosphere depend on the availability of water. Along with other human requirements, a huge amount of water is also used for industrial processes. Among different industries, the textile industries

utilize an enormous amount of water for different activities such as dyeing and washing and similarly this untreated water becomes part of the environment as wastewater (Imran et al., 2015). It is reported that, after dyeing one kilogram (Kg) of fabric, about 40–65 L of wastewater is produced (Balarak et al., 2019). Approximately 20 % of applied quantity of different dyes is lost during the process of washing and dyeing and discharged along with wastewater (Fathima et al., 2018). The major

Abbreviations: BOD, Biological oxygen demand; COD, Chemical oxygen demand; EC, Electrical conductivity; Cu-NPs, Copper nanoparticles; UV-vis, Ultraviolet visible spectroscopy; SEM, Scanning electron microscopy; FTIR, Fourier transform infrared spectroscopy; XRD, X-ray diffraction.

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problem with untreated textile wastewater is its color due to the presence of these dyes in large quantities. It is reported that dyes have many carcinogenic and mutagenic properties. Moreover, several environmental problems such as elevation of various parameters like pH, BOD, suspended bodies, and COD of aquatic bodies are associated with untreated textile effluents (Imran et al., 2015, Subramanian et al., 2022; Bilal et al., 2023). Thus, dyes loaded textile wastewaters limits downstream usage such as drinking, washing, irrigation, and recreation (Imran et al., 2015).

Many physical (adsorption, membrane filtration, coagulation and filtration method), chemical (oxidation like chlorination, ozonation, bleaching and advanced oxidation method with heterogenous and homogenous catalysts) and biological (employing bacteria, fungi, yeast, algae, and plants) methods have been used for the treatment of dyes loaded textile wastewater (Singh et al., 2017). Above mentioned physiochemical methods are associated with many drawbacks such as they are expensive and make use of dangerous chemicals. Furthermore, they are also a source of secondary pollutants such as production of sludge (Bharagava et al., 2020). In contrast, biological methods are environmentally-friendly and even they have ability to completely mineralize many azo dyes. Many reports have described the role of microorganisms in the removal of azo dyes from textile effluents (Husain et al., 2013, Thangaraj et al., 2021). Microorganisms involved in decolorization of azo dyes produce many enzymes such as azo reductase, laccase, peroxidase, MG reductase, and aminopyrine N-demethylase (Souza et al., 2022). However, the rate of biodegradation process depends on many factors including survival activity, and adaptability of microbes with chemical properties of azo dyes and other xenobiotics (Manjarrez Paba et al., 2021). Although biological methods are eco-friendly but the old methods may not be much effective to fulfill the strict standards of water quality and to remove emergent contaminants.

During the past few years, use of nanomaterials has attracted worldwide attention for the degradation of various dyes. They are preferred because of production of biodegradable end products such as aromatic amines which can be further biodegraded easily by microbes (Nazar et al., 2018). Various studies have highlighted the catalytic potential of different metal nanoparticles including silver, titanium, zinc and nickel nanoparticles for the decolorization of different dyes (Edison et al., 2016, Saravanan et al., 2017, Kiran et al., 2020, Mustafa et al., 2023). Among other metal nanoparticles, copper nanoparticles are of excessive interest because of their low cost, availability, and unique properties such as catalytic, antibacterial, biocidal, optical, magnetic and heat transfer abilities (Punniyakotti et al., 2020, Ghaffar et al., 2021). Furthermore, in comparison to silver-nanoparticles, copper nanoparticles (Cu-NPs) have more chemical and physical stability and ease of mixing with polymers (Palani and Elangovan, 2022). The Cu-NPs find their applications in many fields like health (antibacterial, anticancer, dentistry, etc.), food and agriculture (antifungal, nano herbicides, nano-fertilizers, improving texture, etc.), industry (nano composites, cosmetics, nano-pigments, paper coating) and environment (nano remediation, wastewater treatment, biodegradation of polymer, environmental catalysts, etc.) (Ball et al., 2019). For example, Cu-NPs have the ability for dechlorination of dichloromethane (a ground water contaminant) with NaBH_4 by using them as catalyst (Huang et al., 2012). They showed 90 % reduction of dichloromethane within 1 h using NaBH_4 and no degradation was observed without NaBH_4 (Huang et al., 2012).

Different physical and chemical methods like precipitation-pyrolysis method, sonochemistry, cathodic vacuum arc and solvothermal reactions have been reported for physicochemical synthesis of copper-based nanoparticles (Botsa et al., 2019). All such methods make use of the chemicals which make them eco-hazardous and avoid their applications in other fields of clinical, medicine and biology (Akindelu et al., 2020). So, the researchers are more interested toward biological method for nanoparticles synthesis because of the above-mentioned problems that are associated with physicochemical methods. Numerous bacterial

species are reported to synthesize copper nanoparticles like *Salmonella typhimurium* (Ghorbani et al., 2015), *Shewanella oneidensis* (Kimber et al., 2018), *Halomonas elongate* (Rad et al., 2018), *Streptomyces* sp. (Hassan et al., 2019), *Escherichia coli* (Zaynitdinova et al., 2020), *Pseudomonas stutzeri* (Zaynitdinova et al., 2020), *Pseudomonas fluorescens* (Bayat et al., 2021). Similarly, various studies have highlighted the capability of copper nanoparticles in facilitating the photocatalytic degradation of different dyes (Fathima et al., 2018, Noman et al., 2020). For wastewater treatment, Anjum et al. (2019) described the nanotechnology as one of the most advanced process. In this backdrop, following research investigates an innovative and sustainable approach for synthesizing Cu-NPs. The potential of bacterial strain was harnessed with an aim to not only reduce the environmental impact but also enhance its cost effectiveness. For in-depth understanding of Cu-NPs, various cutting-edge techniques such as, UV-Vis spectroscopy, FTIR, SEM, and XRD were used. This study was not only focused on green synthesis but also holds an immense promise in catalytic biodegradation of various azo-dyes. Furthermore, the potential of nanoparticles for the treatment of textile waste water was also explored, thus contributing to environmental conservation. This is the research that promises to not only captivate but also contribute to a greener and more sustainable future.

2. Materials and methods

2.1. Materials

All the reagents or chemicals namely nutrient agar, nutrient broth, and copper sulfate pentahydrate, used for experimental analysis, were procured from Sigma Aldrich or UNI-Chem and used without any purification techniques. The dyes used in this study namely congo red (CR), reactive black-5 (RB-5), methylene blue (MB), reactive red-2 (RR-2), and blue direct (BD) were also purchased from Sigma Aldrich. Moreover, the glassware used during experimental analysis was washed with distilled water and autoclaved.

2.2. Collection of samples for the isolation of bacterial strain

Wastewater samples were collected from Five-star Textile Industry Faisalabad. Bacterial strains were isolated using serial dilution method. Dilutions (10^{-4} to 10^{-6}) were spread on plates of nutrient agar amended with the solution of copper sulfate (1 mM). These plates were incubated for 48 h in incubator at 30 °C. The bacterial colonies having ability to tolerate stress were purified by repeated streaking method on freshly prepared plate and labelled. Among different isolates, strain FB-1 was used for the biosynthesis of nanoparticles.

2.3. Biosynthesis of copper nanoparticles

For the biosynthesis of copper nanoparticles (Cu-NPs), the colony of bacterial strain was inoculated in nutrient broth medium along with two controls (The first control containing only the uninoculated medium and the second control containing the uninoculated medium along with the precursor salt), and incubated under shaking condition at 120 rpm for 48 h at 28 °C. The culture was centrifuged for 10 min at 7000 rpm to remove the pellet for extracellular synthesis of Cu-NPs. The supernatant was then added with 10 mM copper sulfate pentahydrate solution and incubated again under shaking condition for 24 h at 120 rpm. Equal concentration of precursor salt solution was also added to the second control. The change in color from bluish-to-bluish green in comparison to control confirmed the biosynthesis of Cu-NPs. The reaction mixture was again subjected to centrifugation for about 10 min at 7000 rpm at 4 °C to remove the supernatant. The pellet was washed with double distilled water to remove the impurities and placed in an oven to heat dry the nanoparticles. They were then ground into fine powder.

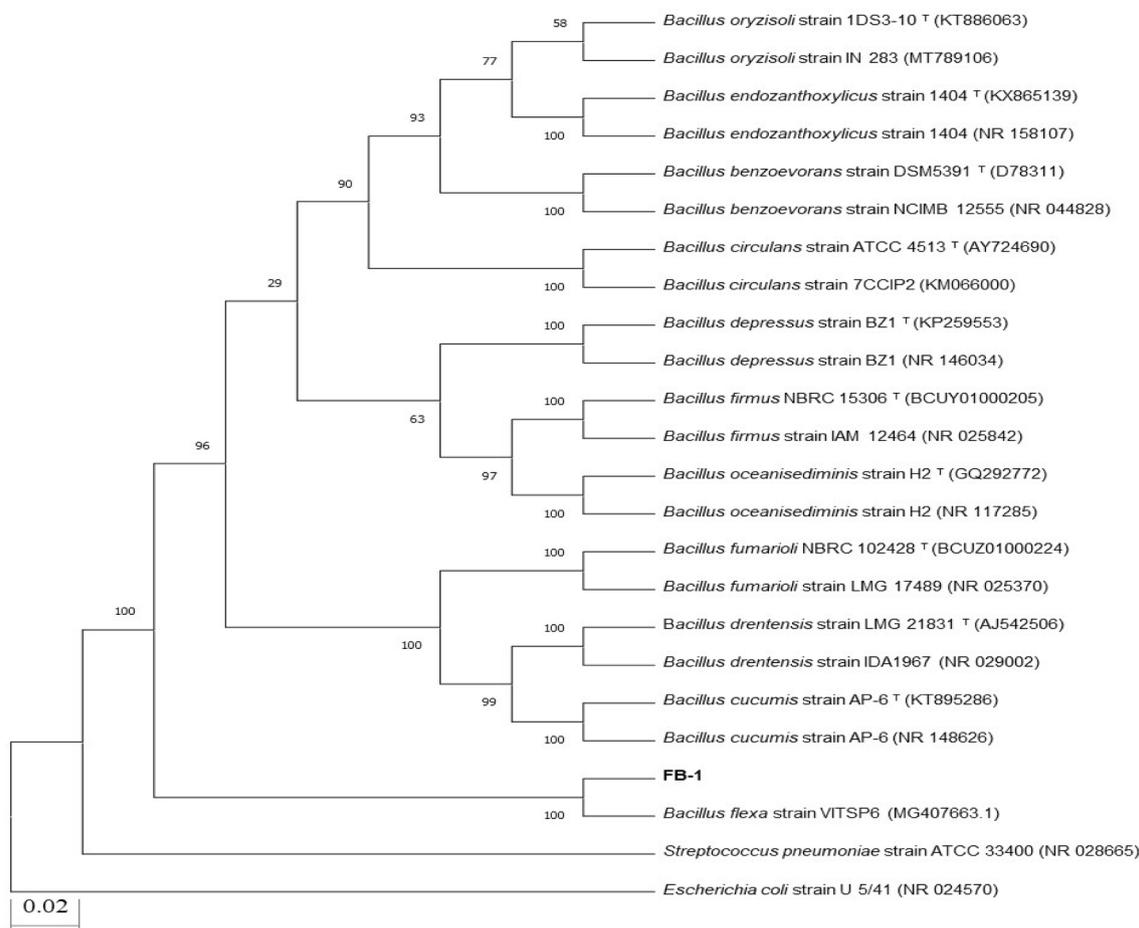


Fig. 1. Neighbor joining phylogenetic tree of *Bacillus* spp. Strain FB-1.

2.4. Molecular characterization and phylogenetic study

The selected bacterial isolate having potential of Cu-NPs synthesis was identified on molecular basis. The 16 S rRNA gene was amplified using forward and reverse primers f-D1 (5'-AGAGTTTGATCCTGGCTCAG-3') and r-D1 (5'-AAGGAGGTGATCCAGCC-3') (Weisburg et al., 1991). The amplified gene products of 16 S rRNA were sequenced commercially by Macrogen, South Korea. The retrieved sequence of 16 S rRNA gene was trimmed analyzed by Sequence Scanner software package and trimmed on both sides to cut the bad sequences. The BLASTn was used for identification of strain. Then the phylogenetic tree was constructed by using NJ (neighbour-joining) method to describe the identity of selected strain.

2.5. Characterization of biosynthesized copper nanoparticles

The bacterial synthesis of Cu-NPs was confirmed using Ultraviolet Visible Spectroscopy (Shimadzu, Kyoto, Japan). It determined the maximum absorption spectra of samples. UV-vis spectroscopy analyzed the reaction mixture at a wavelength ranging from 300 to 800 nm (Noman et al., 2020).

Scanning Electron Microscopy (SEM) was employed to describe the shape and surface texture of biosynthesized Cu-NPs (Instrument: JSM-6490). SEM was conducted by following the method of Tiwari et al. (2016). For SEM analysis, about 10 μ L of homogenous cell free colloidal solution was placed on cover slip made up of glass and dried at 100 $^{\circ}$ C. After that, the glass cover slip was cooled and fixed on aluminum stub using adhesive carbon tape.

The crystalline nature of biosynthesized nanoparticles was deter-

mined using X-ray diffraction analysis. To determine the XRD pattern, the drop was coated on glass substrate and placed in X-ray diffractometer apparatus (JCPDS Card # = 04-0836) which was operated at 40 mA and 45 KV voltage of current with Cu K α rays. The crystalline nature of biosynthesized nanoparticles was analyzed, and Debye-Scherrer's formula, was used for their average size

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

Whereas,

D = "Crystalline size", K = "Sharrer's constant" and its value is "0.9", λ = Wavelength of "x-ray source", β = "Width at 1/2 maximum of reflection peaks" and, θ = "Peak position".

Functional groups and associated proteins that participate in stability of biosynthesized Cu-NPs were determined by Fourier-Transform Infrared Spectroscopy (FT-IR) (FTIR-Bruker TENSOR- 27). The dried powder of biosynthesized Cu-NPs was taken to FTIR measurement and the FTIR spectra were attained. It is useful method to ascertain the various type of chemical bonds in a molecule by making an IR absorption spectrum that functioned as molecular fingerprint. The analysis of FT-IR was conducted in a range of 4000–350 cm^{-1} .

2.6. Catalytic degradation of azo dyes

2.6.1. Optimization of concentration of Cu-NPs for RB-5 decolorization

Various concentrations of Cu-NPs were used to check their potential of decolorization of RB-5. 50 ppm concentration of RB-5 was used for this purpose. Various concentrations of NPs such as (0.05, 0.1, 0.2, 0.5,

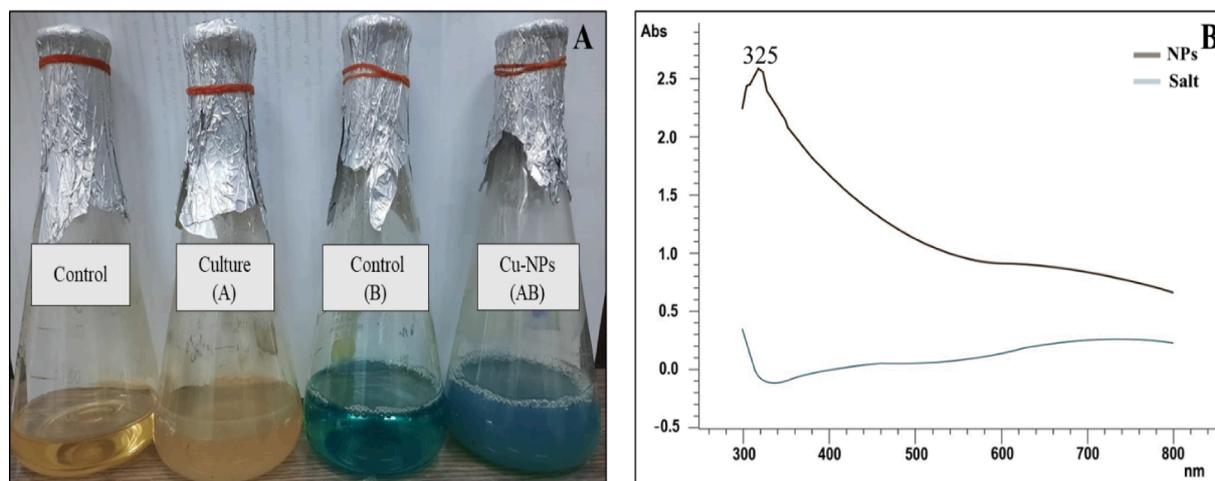


Fig. 2. (A) Confirmation of biosynthesized copper nanoparticles by changing in color and (B) UV-vis spectroscopy. The brown line in UV-Vis spectrum confirmed the synthesis by showing an absorption peak at 325 nm while the blue line indicates the control. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

1, 2, 3, 4, 5 and 10 mg ml⁻¹) were added to the solution of RB-5 and placed in solar irradiation for 4 h after shaking for about 10 min. After each hour, 1 ml of sample was taken from each concentration in a 1.5 ml Eppendorf tube. The samples were centrifuged for about 10 min at 7000 rpm to remove the Cu-NPs. Spectrophotometer was used to check the ability of different concentrations of nanoparticles by measuring the decolorization of RB-5. The optimized concentration (0.5 mg ml⁻¹) of Cu-NPs was used for further analysis.

2.6.2. Decolorization of azo dyes using optimized concentration of NPs

The optimized concentration (0.5 mg ml⁻¹) of nanoparticles was used for the decolorization of various other dyes such as MB, RB-5, BD, CR, MG, and RR-2 dyes at 25 and 50 ppm concentrations. For this purpose, optimized concentration of biosynthesized Cu-NPs (50 mg) was added to the solution (100 ml) of above-mentioned azo dyes. The reaction mixtures were then placed in sun light for incubation for about 4 h. 1 ml of sample was taken from each dye in 1.5 mL Eppendorf tube after specific time interval. The Eppendorf tubes containing samples were centrifuged for 10 min at 7000 rpm to remove the Cu-NPs. Spectrophotometer was used to evaluate the degradation of dyes. The decolorization of dyes namely MB, CR, BD, RR-2, RB-5, and MG was evaluated at 561, 478, 592, 540, 597, and 624 nm, respectively. The % age of dyes degradation was calculated using the formula written below,

$$\% \text{ decolorization} = ((C - S) \times C) / 100$$

Whereas,

- C is control
- S is sample

2.6.3. Treatment of wastewater

Bacterial synthesized Cu-NPs were also used for the treatment of actual wastewater. The sample of wastewater was collected from paharang drain (North-31.526904 and East-73.118483) and Sargodha Road, Faisalabad, Pakistan. It was firstly centrifuged at 10,000 rpm for 5 min to remove the particulate matter. The collected wastewater was spiked with congo red (50 ppm) to have considerable concentration of the dye and color intensity. 50 mg (0.5 mg ml⁻¹) of biosynthesized Cu-NPs were then added to 100 ml of wastewater by keeping the control without nanoparticles to evaluate the treatment of wastewater. Sample was vortexed and incubated in the presence of sunlight for about 7 h. After incubation, the samples were centrifuged for 10 min at 7000 rpm to remove the Cu-NPs. Different parameters like pH, chemical oxygen

demand, electrical conductivity, phosphates, color intensity, and sulphates were checked before and after the treatment following the procedure described by Rump (1999) and Maqbool et al. (2016).

2.7. Statistical analysis

The experimental data of treatment was entered in Microsoft Excel 2016, and percentage decolorization, means, and standard deviation was calculated. For analyzing data, completely randomized design was used. Analysis of variance was calculated by using statistix 8.1 software. Principle component analysis and heat map was also constructed using the R-studio (Version 4.2.2).

3. Results and discussion

3.1. Isolation, molecular characterization and phylogenetic analysis

The selected isolate was purified by repeated streaking method on freshly prepared nutrient agar plates until single colony was obtained. The growth pattern of selected isolate on nutrient agar plate media has been represented in the Figure S1. The phylogenetic analysis of 16 S rRNA gene and BLASTn of isolate determined that the selected isolate belongs to genus *Bacillus*. In Phylogenetic tree (Fig. 1), selected strain clustered with *Bacillus* spp. The 16S rRNA gene sequence of strain (FB-1) was also compared with known sequences in database using BLASTn program of NCBI and the sequence showed more than 99 % resemblance with the bacterial strains belonging to genus *Bacillus*.

3.2. Biosynthesis and characterization of copper nanoparticles

Various microbial isolates have been reported describing the synthesis of Cu-NPs. In this study *Bacillus* spp. Strain FB-1 was used for the extracellular synthesis of Cu-NPs at 10 mM concentration of copper sulphate salt. On addition of salt, the change in color (from bluish to blue greenish) firstly confirmed the biosynthesis process (Fig. 2A). Similar result was reported earlier by Tiwari et al. (2016) who synthesized Cu-NPs using *Bacillus cereus* strain at 10 mM concentration of copper sulfate salt. On the other hand, Noman et al. (2020) also reported the bacterial synthesis of Cu-NPs using *Escherichia* sp. strain SINT7 at 5 mM concentration of precursor salt. The *Bacillus flexus* strain was also used for the synthesis of silver nanoparticles (Priyadarshini et al., 2013). Furthermore, several other nanoparticles such as cadmium sulphide (Singh et al., 2011), iron oxide (Sundaram et al., 2012), titanium dioxide (Khan and Fulekar, 2016) and zinc oxide (Hamk et al., 2023) were synthesized

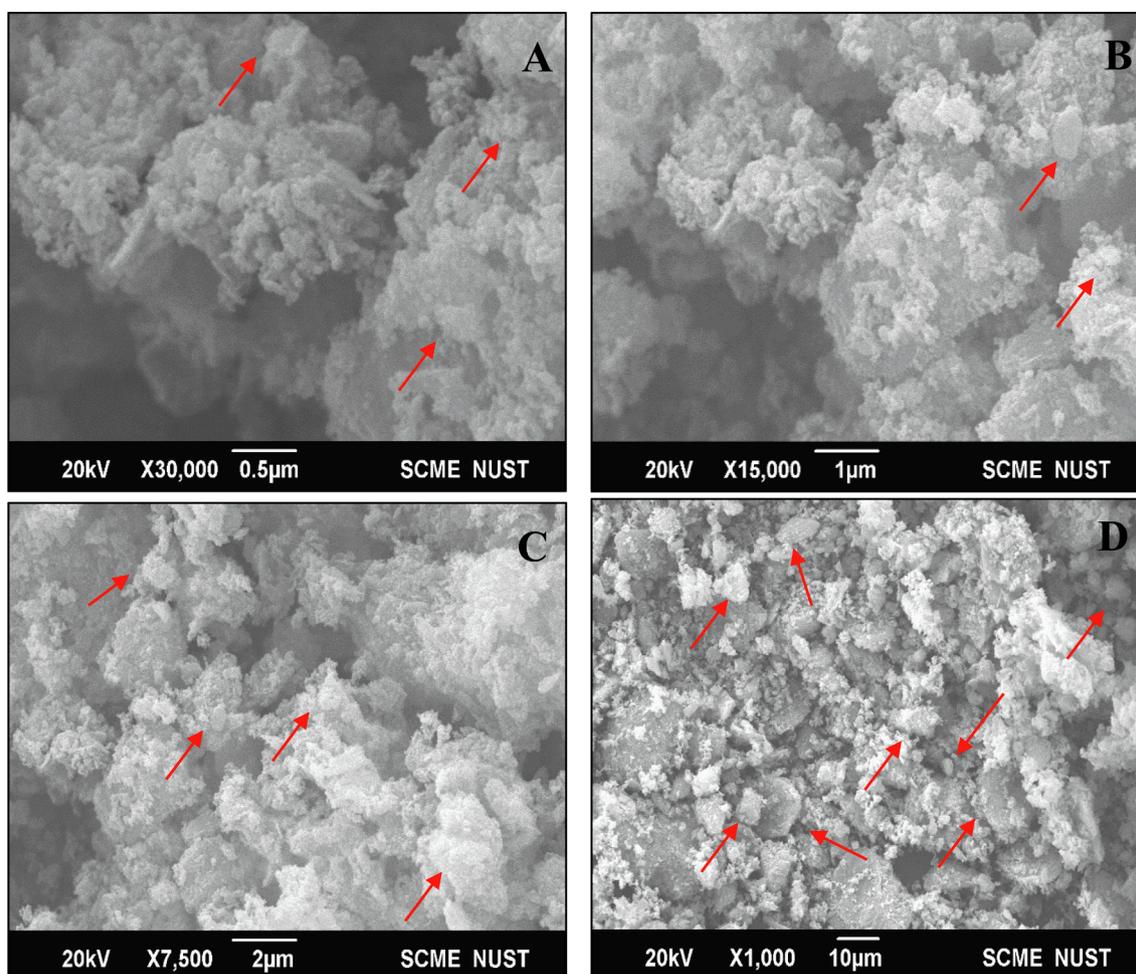


Fig. 3. SEM images of biosynthesized copper nanoparticles under different magnifications such as 0.5 μm (A), 1 μm (B), 2 μm (C), and 10 μm (D).

from *Bacillus* spp. Hence, the bacterial strain *Bacillus* spp. FB-1 is a new addition as a bacterial member for the synthesis of nanoparticle.

The change in color was the first confirmation of biosynthesis of Cu-NPs. The cell free reaction mixture showed absorbance peak at 325 nm using UV–vis spectrophotometer which confirmed the synthesis of Cu-NPs as depicted in Fig. 2B. Similar result was reported by Noman et al. (2020) and concluded that peak in this region confirmed the reduction of precursor salt into nanoparticles. Moreover Singh et al. (2010) and Mamuru et al. (2016) also reported the confirmation peak of Cu-NPs at 310 nm. While Fathima et al. (2018) and Hassan et al. (2019) showed an absorption peak at 560 and 590 nm.

The SEM analysis revealed the morphology of biosynthesized Cu-NPs. The SEM image of the biosynthesized Cu-NPs (Fig. 3) indicated that the sample contained copper nanoclusters with irregular spherical shape and present in an agglomerated form. The analysis of SEM images revealed that the particles size was in range of 17–34 nm. Various reports described the different shapes of Cu-NPs such as cubical (Kuo et al., 2007), rhombic dodecahedral (Liang et al., 2009), 18-facet polyhedral (Lin et al., 2010), octahedral (Xu et al., 2011) and spherical (Noman et al., 2020), Bukhari et al., 2021).

The XRD analysis of biosynthesized Cu-NPs showed various diffraction peaks at 44.5° , 51.5° , 74.75° as depicted in Fig. 4A. The described peaks at 44.5° , 51.5° , 74.75° were assigned to (111, 200 and 220) planes of copper. The peaks at above mentioned planes confirmed the crystalline nature of Cu-NPs. There are also some extra peaks that might be due to presence of some cupric oxide or copper oxide nanoparticles. The size of nanoparticles was less than 27.6 nm calculated by Debye-Scherrer's formula. Tiwari et al. (2016) reported the similar results and showed

diffraction peaks of Cu-NPs at 111, 200 and 220 synthesizing from *Bacillus cereus* spp. They also reported some extra peaks of copper oxide nanoparticles. Similarly, Noman et al. (2020) and Lv et al. (2018) also reported the similar results showing diffraction peaks at 111, 200, and 220 with a size of particles about 28.55 and 10–16 nm.

FTIR spectrum of biosynthesized Cu-NPs produced from *Bacillus* sp. FB-1 showed peaks of absorption at 3388.72, 2928.94, 1645.23, 1403.43, 1066.48, and 595.02 cm^{-1} as shown in Fig. 4B. The peaks at 3388.72 and 2928.94 cm^{-1} were due to presence of OH group of alcohol and C–H stretching. The peak at 1645.23 cm^{-1} was due to C=C stretching. Whereas peaks at 1403.43 and 1066.48 cm^{-1} were due to bending of C–H and stretching of C–O group. The presence of bands at 595.02 cm^{-1} showed probabilities of existence of copper or copper oxide nanoparticles (CuO-NPs). The FTIR spectrum of biosynthesized Cu-NPs confirmed the presence of alcoholic group and proteins. Coating of protein around copper nanoparticles enhanced the long-term stability of particles (Valodkar et al., 2011). Whereas, Noman et al. (2020) reported the various absorption peaks at 3381.79, 2957.81, 1656.20, 1451.87, 1398.87 and 1111.12 cm^{-1} . The peaks at 3381.79 and 2957.81 cm^{-1} were due to presence of OH group of alcohol and stretching of C–H bond. While Nieto-Maldonado et al., (2022) reported the plant based synthesis of copper nanoparticles and showed various absorption peaks at 3271, 2931, 1606, and 1029 cm^{-1} . The peaks at 3271 and 2931 cm^{-1} were due to presence of O–H and C–H groups. On the other hand, the peaks at 1606 and 1029 cm^{-1} were due to C=O and C–O groups.

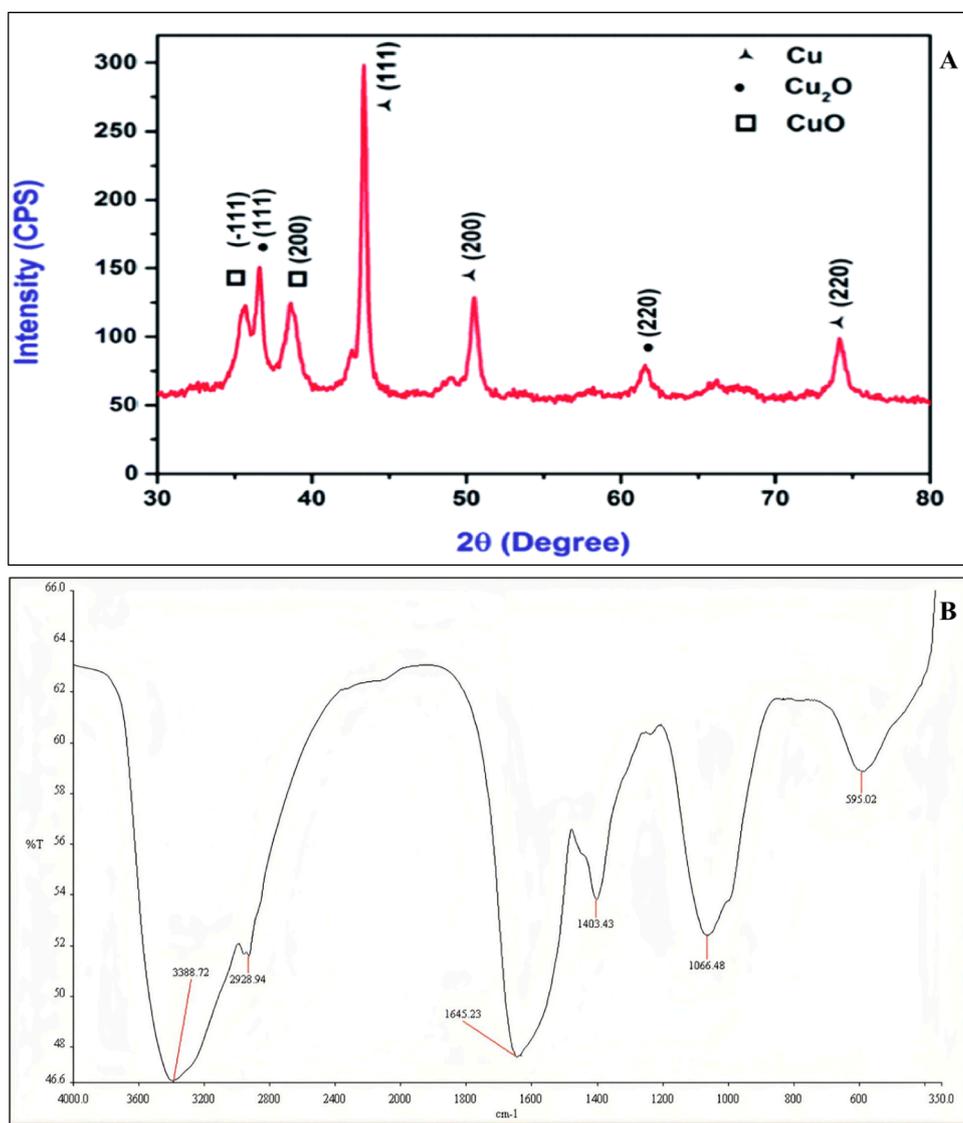


Fig. 4. XRD (A) and FTIR (B) spectrum of biologically synthesized copper nanoparticles using *Bacillus* spp.

3.3. Photocatalytic degradation of azo dyes

3.3.1. Optimization of concentration of Cu-NPs for RB-5 decolorization

In the current work, different concentrations of biosynthesized Cu-NPs were used for the degradation of RB-5. After solar incubation of 4 h, decolorization of RB-5 with 0.05, 0.1, 0.2, 0.5, 1, 2, 3, 4, 5, and 10 mg ml⁻¹ concentration of biosynthesized Cu-NPs was about 84.3 ± 0.3 %, 87.0 ± 0.5 %, 87.2 ± 0.5 %, 86.9 ± 1.4 %, 90.1 ± 1.8 %, 89.0 ± 1.6 %, 89.7 ± 2.6 %, 90.1 ± 1.2 %, 88.0 ± 3.1 %, and 88.5 ± 3.4 %, respectively (Fig. 5). From the rate of decolorization, it was clearly proved that each concentration shows maximum degradation of RB-5. However, it is noteworthy that, after 1 h incubation period, 24.9 ± 2.9 %, 45.0 ± 0.3 %, 60.4 ± 1.6 %, 80.5 ± 0.8 %, 80.8 ± 1.2 %, 80.8 ± 0.2 %, 81.9 ± 0.7 %, 80.9 ± 1.3 %, 81.8 ± 2.5 % and 75.5 ± 2.1 % decolorization of RB-5 was observed using 0.05, 0.1, 0.2, 0.5, 1, 2, 3, 4, 5, and 10 mg ml⁻¹ concentration of Cu-NPs. So, 0.5 mg ml⁻¹ of biosynthesized Cu-NPs was selected for the decolorization of other dyes as shown in Fig. 8. While Noman et al. (2020) used 1 mg ml⁻¹ of Cu-NPs for the decolorization of various azo dyes. Furthermore Ahmad et al. (2018) decolorized the bromophenol blue (BPB) dye using 0.25 mg ml⁻¹ of green synthesized Cu-NPs.

Histogram correlation analysis was carried out for the selection of best concentration of nanoparticles for the decolorization of dyes. In a

histogram (Fig. 6A) red color showed non-significant while pink, grey and blue colors showed significant results. After analyzing the histogram, it was observed that among all concentrations, 0.5 mg ml⁻¹ showed exceptional results. When compared to 0.5 mg ml⁻¹ concentration, lower concentrations (0.05–0.2 mg ml⁻¹) showed significant decolorization but not effective as 0.5 mg ml⁻¹ concentration. On the other hand, higher concentrations (1–10 mg ml⁻¹) did not showed a proportional increase in decolorization potential, especially the minimum increase was observed after the 3 h. While, 0.5 mg ml⁻¹ showed a higher initial decolorization rate and maintained throughout the time, thus providing an optimum balance between effectiveness and nanoparticles concentrations. So, the 0.5 mg ml⁻¹ was selected that not only conserve the resources but also go with environmental and economic considerations, potentially reducing the ecological footprint and material costs associated with the process of decolorization of dyes.

The loading plots of PCA to check the effect of different concentrations of nanoparticles on rate of decolorization have been presented in Fig. 6B. In the whole database, Dim-1 and Dim-2 showed maximum contribution and occupy more than 98.5 % of all databases, among which Dim-1 showed 94.1 % and Dim-2 showed about 4.4 %. It was observed that concentration of nanoparticles showed a significant effect on rate of decolorization.

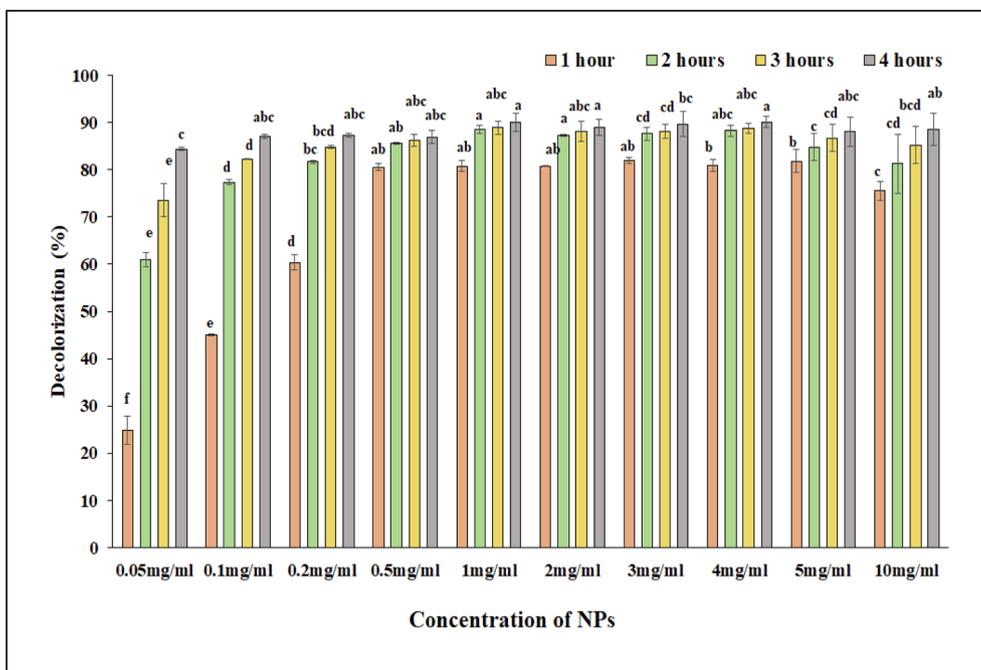


Fig. 5. Photocatalytic degradation of RB-5 with different concentrations of biosynthesized copper nanoparticles (0.5, 0.1, 0.2, 0.5, 1, 2, 3, 4, 5, and 10 mg/ml) at 50 ppm of dye concentration, respectively.

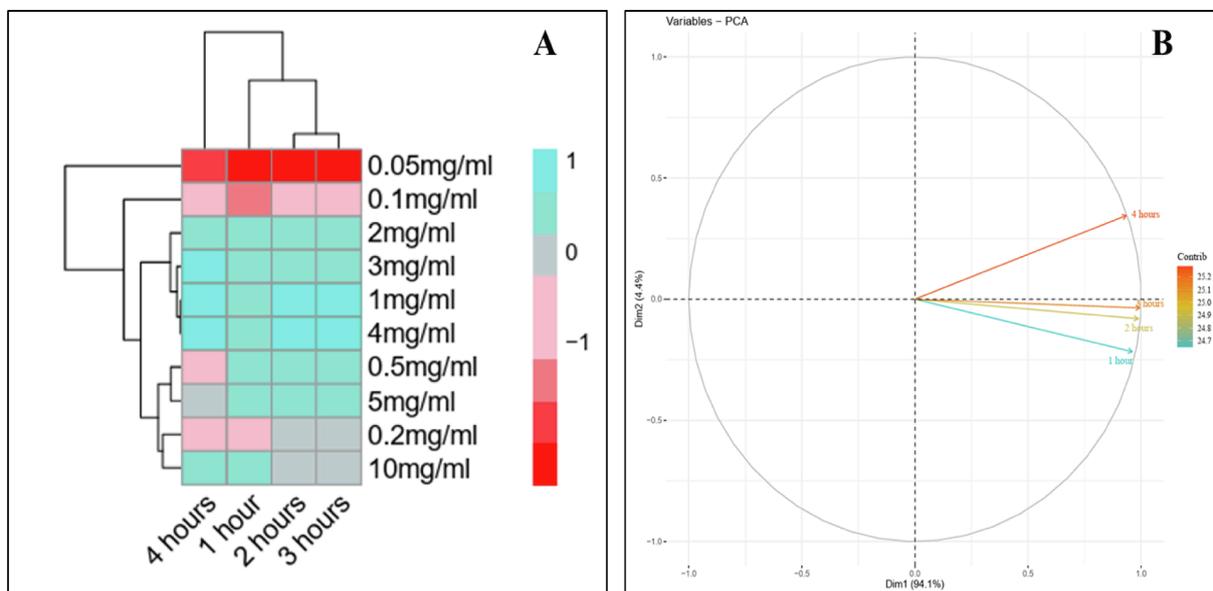


Fig. 6. Heat map (A) and Principle component analysis (B) using different concentrations of nanoparticle for the decolorization of RB-5 dye under solar irradiation of 4 h.

3.3.2. Decolorization of azo dyes using optimized concentration of NPs

After optimization, 0.5 mg ml⁻¹ concentration of biosynthesized Cu-NPs was used for the decolorization of various other azo dyes. The Cu-NPs were observed to carry out an effective decolorization of different azo dyes (Fig. 7). The results indicated that, just after solar irradiation of 1/2 h, Cu-NPs decolorized about 84.5 ± 1.1 %, 64.8 ± 3.9 %, 38.7 ± 4.3 %, 46.9 ± 2.2 %, 20.4 ± 3.8 % and 30.7 ± 2.1 % of RB-5, CR, MG, MB, RR-2, and BD. However, 71.9 ± 1.9 %, 30.5 ± 1.4 %, 61.9 ± 3.9 %, 8.1 ± 0.5 %, 15.7 ± 2.2 % and 15.9 ± 1.6 % of RB-5, CR, MG, MB, RR-2, and BD were decolorized at 50 ppm concentration of dyes just after 1/2 h. However, after 1 h of solar incubation, decolorization of RB-5, CR, MG, MB, RR-2, and BD was increased to 86.3 ± 0.7 %, 90.3 ± 2.4 %,

54.2 ± 9.4 %, 58.9 ± 3.5 %, 36.7 ± 3.3 % and 45.8 ± 2.1 % at 25 ppm concentration. Similarly, at 50 ppm concentration, the decolorization of RB-5, CR, MG, MB, RR-2, and BD was also increased to 84.2 ± 0.6 %, 71.6 ± 2.5 %, 71.6 ± 3.9 %, 46.1 ± 2.8 %, 26.8 ± 1.6 % and 48.5 ± 5.5 %, after 1 h of solar incubation. It is noteworthy that the decolorization of the above-mentioned dyes was more at 25 ppm (Fig. 7A) concentration of dyes than 50 ppm (Fig. 7B) concentration. Finally after 4 h of solar irradiation, the copper nanoparticles decolorized about 96.8 ± 1.0 % of RB-5, 92.2 ± 2.2 % of CR, 96.9 ± 1.1 % of MG, 72.5 ± 0.6 % of MB, 58.5 ± 1.2 % of RR-2 and 78.7 ± 2.7 % of BD at 25 ppm concentration and 88.2 ± 0.2 % of RB-5, 88.5 ± 1.9 % of CR, 90.4 ± 1.8 % of MG, 64.8 ± 1.0 % of MB, 46.8 ± 1.6 % of RR-2, and 67.3 ± 2.9 % of BD at 50 ppm

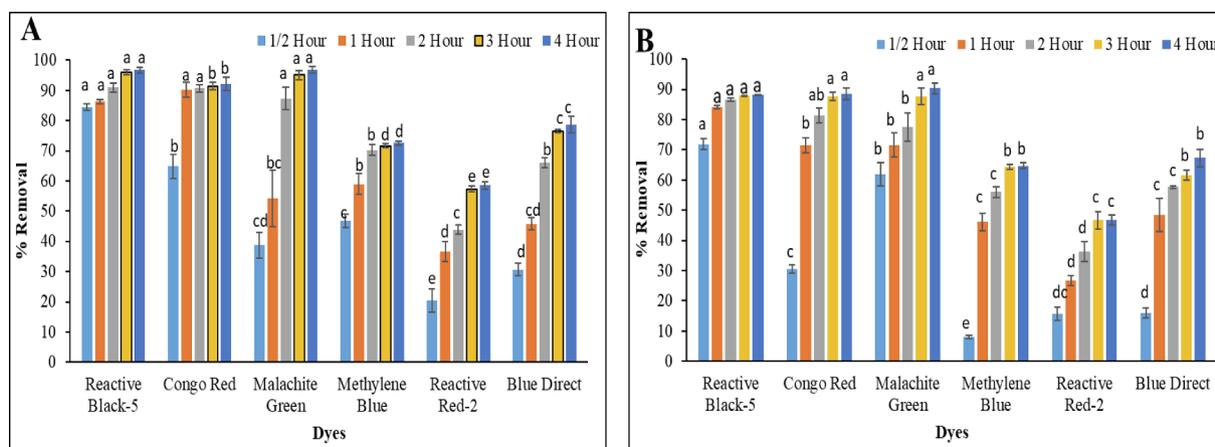


Fig. 7. Decolorization of RB-5, CR, MG, MB, RR-2 and BD under solar irradiation of 4 h at 25 (A) and 50 (B) ppm concentration.

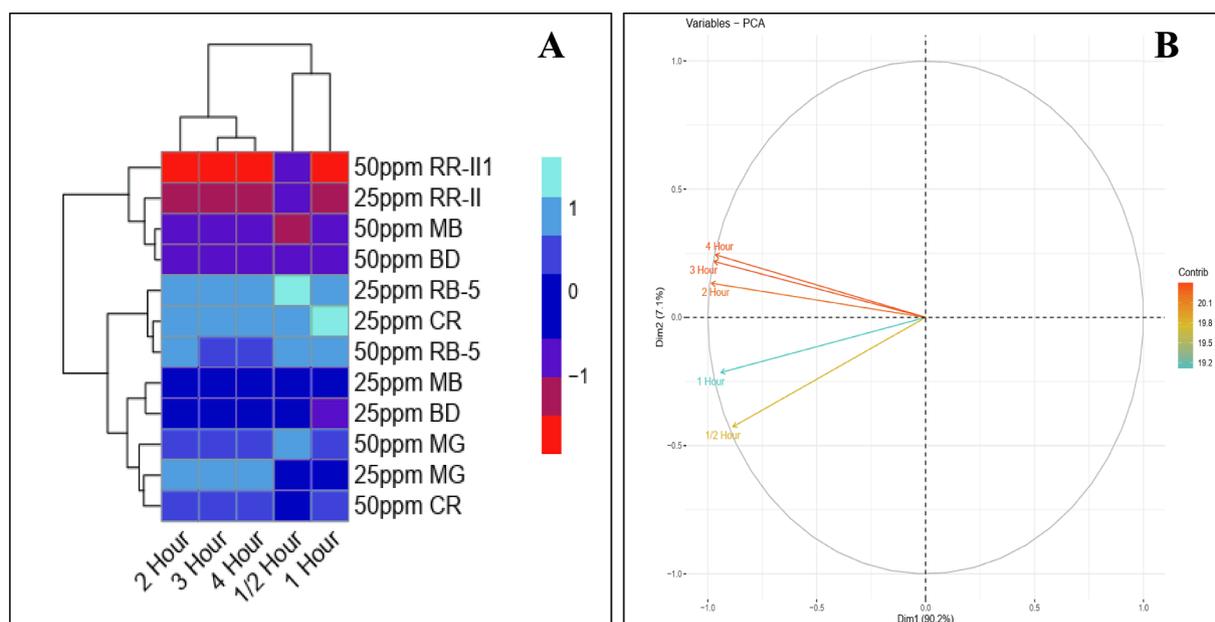


Fig. 8. Heat map (A) and Principle component analysis (B) using different concentrations of dyes under solar irradiation of 4 h.

concentration. The difference in catalytic potential of biosynthesized Cu-NPs for different dyes might be due to change in activity in aqueous media, structure of dye, and difference in intensity of solar radiation (Noman et al., 2020). The visual representation of decolorization of above-mentioned dyes has been shown in Figure S2. Dyes were decolorized after the addition of biosynthesized Cu-NPs and was proved from the decrease in absorption values at λ_{max} of each dye. While, (Noman et al., 2020) reported the 97.07 %, 83.61 %, 88.42 %, and 90.55 % decolorization of 25 ppm concentration of congo red, reactive black-5, direct blue-1, and malachite green respectively. On the other hand, Ghaffar et al. (2021) reported the 73.5 % decolorization of Disperse yellow-125 using 0.01 % concentration of dye and 0.05 % of green synthesized Cu-NPs.

While, the photocatalytic activity of Cu-NPs for the degradation of various azo dyes such as, congo red, methyl red, methylene blue (Fathima et al., 2018), and methyl orange (Li et al., 2015) was also reported (Al-Hakkani, 2020). Moreover, the catalytic potential of biosynthesized nanoparticles can be used for the removal of pollutants present in actual wastewater.

The heat map analysis was also carried out to check the effect of different concentration of dyes over time. In a histogram (Fig. 8A) red

color showed non-significant while pink, grey and blue colors showed significant results. After analysis it was observed that for both concentrations (25 and 50 ppm), the decolorization potential of biosynthesized Cu-NPs generally increased over time, with some variances among dyes. At 25 ppm concentration, RB-5, CR, and MG showed high decolorization potential, reaching over 90 % after 4 h. The decolorization of MB and BD also increased over time, while RR-II has the least decolorization potential. On the other hand at 50 ppm concentration, RB-5 showed a consistent increase in decolorization potential, but slightly lower than the 25 ppm concentration. While decolorization potential of CR, MG, and BD showed significant improvements over time, and MB and RR-II have moderate to low decolorization potential, thus indicating that higher concentrations do not lead to better decolorization for these dyes.

Similarly PCA was also used to check the effect of different concentrations of dyes on rate of decolorization (Fig. 8B). In the whole database, Dim-1 and Dim-2 showed maximum contribution and occupy more than 97.3 % of all databases, among which Dim-1 showed 90.2 % and Dim-2 showed about 7.1 %. It was observed that concentrations of dyes showed significant effect on rate of decolorization over time.

Table 1Different parameters before and after the treatment of wastewater with the help of copper nanoparticles synthesized from *Bacillus* sp.

Wastewater treatment	pH	EC (dS m^{-1})	TDS (mgL^{-1})	COD (mgL^{-1})	Sulphates (mgL^{-1})	Phosphates (mgL^{-1})	Color intensity (nm)
Pre-Treatment	8.6	7.31	1940	1644	182.34	27.64	0.4
Post-Treatment	6.9	2.98	821	992	103.65	13.96	0.02
% Removal	–	–	57.68	39.66	43.16	49.49	95.97

3.3.3. Treatment of textile wastewater

The optimized concentration of Cu-NPs was also used for the treatment of textile wastewater added with CR dye (50 ppm). Before any treatment, the wastewater shows high values for various parameters such as sulphates, phosphates, pH, EC, TDS, COD, and color intensity. Whereas the treated wastewater showed significant reduction in all above-mentioned parameters (Table 1). Few studies already reported the treatment of wastewater via metal nanoparticles (Bibi et al., 2019, Dlamini et al., 2019, Al-Hakkani, 2020; Noman et al., 2020, Siddique et al., 2021). The results of present study showed that the pH of the wastewater was reduced from 8.6 to 6.9 after the treatment as compared to non-treated wastewater. The EC of treated wastewater was also reduced by 59.23 %. Whereas, other parameters such as TDS, COD, sulfates, phosphates, and color intensity were also reduced by 57.68 %, 39.66 %, 43.16 %, 49.49 %, and 95.97 % in comparison to non-treated wastewater as depicted in Table 1. Few previous studies have also reported the treatment of wastewaters by exploiting the potential of different types of nanoparticle (Bibi et al., 2019, Noman et al., 2020, Siddique et al., 2021; Rasool et al., 2023; Pham-Khanh et al., 2023). For example, the findings of Siddique et al., (2021) reported the reduction in various parameters such as pH, EC, TDS, COD, and color removal with the help of biologically and chemically synthesized zinc oxide nanoparticles. Furthermore, Fouda et al. (2022) also reported the treatment of tanning wastewater using zinc oxide nanoparticles. They reported the 96.5, 88.8, 88.5, and 91.5 % reduction in TSS, COD, BOD, and conductivity of wastewater treated with zinc oxide nanoparticle. Moreover, zinc oxide nanoparticles also lowered the Cr (VI) level from wastewater by 93.4 % (Fouda et al., 2022). On the other hand, Noman et al. (2020) also treated the textile wastewater with the help of Cu-NPs and discussed the decrease in different parameters like pH, turbidity, COD, TDS, TSS, hardness, chloride, and sulphates before and after the treatment. The reduction efficiency of Cu-NPs from wastewater can be attributed to functional groups, surface structure and chemical components that are attached to them (Dlamini et al., 2019; Al-Hakkani, 2020). So, the above-mentioned results suggested that biosynthesized Cu-NPs might serve as an excellent photocatalyst for the treatment of textile wastewater loaded with dyes.

4. Conclusion

Based on the findings of the current study, it might be concluded that biosynthesized Cu-NPs might serve as potential agent not only for photocatalytic degradation of different textile azo dyes but also for treatment of actual wastewaters by lowering sulphates, phosphates, color intensity, pH, EC, TDS, and COD. Therefore, the current study emphasizes on photocatalytic application of biosynthesized Cu-NPs in the decolorization of azo dyes on commercial level.

Ethics approval

This article does not contain studies with human participants or vertebrates performed by any of the authors.

Consent for publication

All authors give consent for publication of this original article.

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CRedit authorship contribution statement

Fatima Batool: Writing – original draft, Formal analysis, Data curation. **Muhammad Shahid:** Writing – review & editing, Formal analysis, Conceptualization. **Faisal Mahmood:** Writing – review & editing, Resources, Methodology, Formal analysis. **Tanvir Shahzad:** Writing – review & editing, Methodology, Formal analysis. **Farrukh Azeem:** Writing – review & editing, Methodology, Investigation, Conceptualization. **Sabir Hussain:** Writing – review & editing, Supervision, Methodology, Funding acquisition, Formal analysis, Conceptualization. **Tahani Saad Algarni:** Writing – review & editing, Conceptualization. **Mohamed S. Elshikh:** Writing – review & editing, Funding acquisition, Data curation, Conceptualization. **Wed A. Al. Onazi:** Writing – review & editing, Conceptualization. **Sadia Mustafa:** Writing – review & editing, Resources, Data curation.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

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