**ORIGINAL ARTICLE**



# *Taxus wallichiana* **leaf extract‑mediated microwave‑assisted one‑pot biosynthesis of MgO NPs for biomedical and photocatalytic applications**

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# **Abstract**

Potentially useful MgO NPs were synthesized via rapid eco-friendly one-pot microwave-assisted green techniques using *Taxus wallichiana* leaf extract. The development, morphology, and other physicochemical properties of nanoparticles were further analyzed by UV visible spectrometer, FTIR, XRD, SEM, and EDX analysis. The developed nanoparticles were uniformly distributed having particle sizes less than 30 nm. The XRD and SEM analysis of the developed nanoparticles reveals that it is crystalline having almost spherical. Phenol red and rhodamine B are used as potential organic pollutants for the evaluation of the photocatalytic activity of the developed nanoparticles. For the screening of the antibacterial efficiency of the biogenic nanoparticles, fve diferent bacterial pathogens were used. The nanoparticles exhibited signifcant antibacterial potential against the *Bacillus* spp., *Klebsiella pneumonia*, and *Staphylococcus aureus* with a zone of inhibition 23 mm, 20 mm, and 21 mm. The nanoparticles also show moderate antibacterial activity against *Escherichia coli* and *Pseudomonas* spp. with the zone of inhibition 14 mm and 16 mm. Furthermore, the antioxidant potential of the developed nanoparticles was investigated by using a DPPH radical scavenging assay. The present research showed that the developed nanoparticles were the potential low-cost and efective photocatalyst in the treatment of wastewater and also act as potential antibacterial and antioxidants. The use of *Taxus wallichiana* leaf extract in the fabrication of MgO nanoparticles was the frst time reported in the present research work.

**Keywords** Green synthesis · Photocatalytic activity · Antibacterial activity · Magnesium oxide nanoparticles

# **1 Introduction**

Nanotechnology is a transdisciplinary area of science and technology that is involved in the synthesis application and handling of material with a size of less than 100 nm. During the last few decades, nanotechnology has become an emerging feld of research interest [[1\]](#page-8-0). Metal and metal oxide nanoparticles have captivated scrupulous research interest due to their extensive aptness in the diverse feld of applications like the manufacturing of optical devices,

sensors, pharmaceutical applications, and the feld of biotechnology  $[2, 3]$  $[2, 3]$  $[2, 3]$ . The application of nanomaterials in various felds increases extensively and it received much priority as compared to the bulk material due to their large surface area [[4\]](#page-8-3). The nanomaterials exhibit unique physical and chemical properties and the prominent reason behind their unique properties is their very high surface-to-volume ratio. Another important parameter that makes nanomaterials more useful as compared to bulk materials is their discrete energy levels. The bulk material has a continuous energy level [\[5](#page-8-4)]. The development of novel and sustainable methods for the manufacturing of inorganic metal oxide nanoparticles is a great dynamic area of research. Methods involved in the fabrication of nanomaterial are largely classifed into two main categories, physical and chemical methods, but these methods are not very suitable for the synthesis of the materials utilized in the feld of biological and medicinal applications [\[6](#page-8-5), [7](#page-8-6)]. Recently, green technology has drawn more attention to the development of potentially useful



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ceramic nanoparticles during the last few decades due to certain advantages over the available ancient methods [\[8](#page-8-7)].

Green technology uses various kind of plant material and microorganism in the fabrication of diferent kind of potentially useful nanomaterials and the use of these materials significantly reduce the use of harmful toxic chemical [\[9](#page-8-8)]. Hence, green route for the fabrication of nanoparticles is one of the most convenient and eco-friendly approaches for the development of nanoparticles. Diferent phytochemicals like polyphenols, favonoids, vitamins, amino acids, and polysaccharides play a prominent role in the development of nanoparticles via the green route. These phytochemicals have remarkable medicinal value and have eco-friendly behavior. They are good replacements for harmful reducing agents like formaldehyde and sodium borohydride. There is an immense scope to develop new green technology methods for the development of diferent kinds of nanomaterials having adequate antibacterial antifungal and photocatalytic activity [[10–](#page-8-9)[12\]](#page-8-10). The use of the green synthesis method in the manufacturing of nanomaterials enhances the biomedical application of these materials [[13\]](#page-8-11).

The phytoconstituents play a prominent role in the development of nanomaterials. These phytoconstituents not only play as a capping, stabilizing, and reducing agent but also play a crucial role in the modifcation of the shape and size of the nanoparticles [[14\]](#page-8-12).

Clean and safe water is one of the most essential parts of the entire living organism. Water contamination of the present water resources has become one of the most serious issues due to rapid industrialization. Heavy metal ions, dyes, and residues of pharmaceutical and agricultural waste are the major sources of water contamination [[15](#page-8-13)]. Nanomaterials especially metal oxide nanoparticles receive much attention in the treatment waster by the advanced oxidation process. In the advanced oxidation process, highly reactive reaction intermediates such as hydroxyl free radical play a prominent role in the treatment of wastewater and the metal oxide nanoparticles can efficiently produce these reactive species on exposure to solar and UV light [[16\]](#page-8-14).

The ceramic nanoparticles have noticeable optical, physical, chemical, mechanical, and thermo-mechanical properties due to their very high surface-to-volume ratio. Among them, particularly magnesium oxide nanoparticles have notable properties and show excellent photocatalytic biomedical and pharmaceutical applications. It acts as an efficient photocatalyst in the treatment of wastewater due to its wide band energy gap, high thermal conductivity, and good mechanical strength. Tremendous application of magnesium oxide nanoparticles in the feld of sensing, wastewater treatment, as an antimicrobial, in the electronics in clothing, and photography increased its market value enormously [[17\]](#page-8-15).

The use of microwaves in the feld of synthesis of materials has been increasing rapidly due to their rapid heating and consequently reduces the time of synthesis by increasing the rate of reaction  $[18]$ . Not very much significant work has been reported on the use of microwaves in the development of magnesium oxide nanoparticles via the green route. Hence, in this manuscript, we highlight a simple low-cost eco-friendly robust method for the fabrication of MgO NPs by utilizing the aqueous leaf extract of the *Taxus wallichiana.*

# **2 Experimental**

# **2.1 Materials**

The materials used in this study are ethanol (99%), magnesium nitrate (99%), NaOH (99%), phenol red (90%), and rhodamine B (90%). All these chemicals are purchased from the Sisco Research Laboratory Pvt., Ltd. The plant materials are collected from diferent parts of Uttarakhand, India. The plant materials are authenticated by the Botany Department of the FRI Dehradun. The bacterial strains used in this study are received from the Microbiology Department of the university.

# **2.2 Isolation of leaf extract**

The isolated leaf was washed properly with double distilled and it was kept in the room for 4–5 days to remove the moisture. The dried leaf is cut into fne pieces and then converted into fne powder by using an electric blender. Twenty-fve milligrams of fne powder of leaf along with 250 mL of ethanol as a solvent was placed into the Soxhlet extraction assembly. The process of extraction is completed in approximately 4–5 h. When the extraction was completed the isolated leaf extract was stored in an airtight container [\[19](#page-8-17)].

## **2.3 Microwave‑assisted green synthesis of MgO NPs**

For the microwave-assisted green synthesis of MgO NPs, 10 mL isolated leaf extract of *Taxus wallichiana* was mixed in the 50 mL (0.2 M) of freshly prepared Mg (NO<sub>3</sub>)<sub>2</sub> 6H<sub>2</sub>O. The prepared mixture was now subjected to microwave irradiation for about 40–50 s. The rapid change from colorless to brown of the reaction mixture was observed. The color change initially indicates the development of MgO NPs. The developed nanoparticles were collected by centrifugation of the test solution at 20,000 rpm for about 30 min  $[20]$  $[20]$ .

# **2.4 Characterization**

The UV spectroscopic analysis of the developed nanoparticles was performed by using UV visible spectrophotometer (Systonics double-beam spectrophotometer) in the range of 200–800 nm. To appraise the presence of diferent functional groups on the surface of the developed nanoparticles as reducing and stabilizing agents, FTIR analysis was performed by using Thermo Scientifc Nicolet Summit LITE iD1 FTIR. The crystallographic behavior of the developed nanoparticles was explored by using a Rigaku X-ray diffractometer in this analysis; the wavelength of the radiation is 1.5406 Å and operated at 40 kV and 40 mA  $(2\theta = 10-80)$ . The particle size distribution and the morphological behavior of the developed nanoparticles were estimated by using FE SEM (Zeiss evo-18). The metallic compositions of the developed nanoparticles were examined by using (EDX) energy-dispersive X-ray analyzer.

#### **2.5 Photocatalytic activity**

The photocatalytic potential of the developed nanoparticles were examined against the phenol red dye and rhodamine B dye. The 0.25 g of the biologically synthesized MgO NPs was dispersed in the 100 mL aqueous solution of the phenol red and rhodamine B dye. The prepared dye solution was stirred for about 18 h in the dark to establish adsorption and desorption equilibrium between the dye and the prepared nanoparticles. Now the solution was placed in the solar light and UV light (310 nm) and observed any color variation. The photocatalytic potential of the fabricated nanoparticles was evaluated by measuring the absorbance of the centrifuged aliquot (2 mL) of the test solution at a regular interval of 10 min. The maximum absorbance  $(\lambda_{\text{max}})$  of phenol red and rhodamine B dye was measured by UV spectrophotometer. The maximum absorbance for phenol red and rhodamine was observed at 560 nm and 530 nm, respectively [\[21](#page-8-19)].

## **2.6 Antibacterial study**

The antibacterial efficacy of the biologically synthesized MgO NPs was examined by a well-difusion method. For an antibacterial study, the bacterial microorganisms *Bacillus* spp., *Pseudomonas*, *Klebsiella pneumonia*, *Staphylococcus aureus*, and *Escherichia coli* were obtained from the Microbiology Department of the university. The *E coli* is a rod-shaped gram-positive human pathogenic bacteria. The *Bacillus* spp. and *Staphylococcus aureus* are gram-positive bacteria. The *Pseudomonas* and *Klebsiella pneumonia* (anaerobic, rod-shaped bacteria that causes nosocomial infection) are also gram-negative human pathogenic bacteria. The bacterial culture spread on the Mueller–Hinton agar medium at 35 °C. After the solidifcation of the agar medium, 6-mm-diameter four wells were formed by punching with a borer. In the three wells, 30 µL of the diferent concentrations (20%, 40%, and 60%) of green synthesized MgO NPs were inoculated with the help of a micropipette, and in one well gentamycin (50 µg/L) was inoculated as a positive control. The developed plates were incubated at 37 °C for about 24 h and the diameter of the zone of inhibition was measured in mm. The diameter of the zone of inhibition was measured by using the transparent ruler from the back of the bacterial plate [[22](#page-8-20)].

#### **2.7 Antioxidant study**

The antioxidant activity of the developed nanoparticles was investigated by utilizing the 2,2-diphenyl picrylhydrazyl (DPPH) radical scavenging assay. To perform the antioxidant activity frst prepare the diferent concentration (20, 30, 40, 50, 60, 70 μg/mL) of the MgO nanoparticles. Now, 1 mL (0.1 mM) of the DDPH solution was mixed in to test the solution of the MgO NPs, and the test tube was kept in the dark for about 30 min. After 30-min incubation, the absorbance of the test solution was evaluated at 517 nm. Ascorbic acid is used as a reference compound in this study [\[23\]](#page-8-21). The following equation  $(Eq. 1)$  $(Eq. 1)$  $(Eq. 1)$  was used in the evaluation of the radical scavenging activity of the developed MgO NPs.

<span id="page-2-0"></span>% inhibition = 
$$
\frac{A_0 - A_t}{A_0} \times 100
$$
 (1)

where  $A_0$  represents the absorbance of control and  $A_t$  represents the absorbance of the sample.

# **3 Results and discussion**

# **3.1 Characterization**

The UV visible spectral analysis is one of the simplest methods to evaluate the optical properties of microwaveassisted biologically developed MgO NPs. The UV spectra of the developed MgO NPs can be examined in the range of 200–800 nm. The fgure shows a strong absorption peak at 330 nm [\[24,](#page-8-22) [25](#page-8-23)]. The appearance of an absorption peak at 330 nm indicates the formation of MgO nanoparticles (Fig. [1\)](#page-3-0).

# **3.2 Characterization of green synthesized MgO NPs by XRD analysis**

The XRD analysis of the developed MgO NPs is depicted in Fig. [2.](#page-3-1) The developed nanoparticles were found to be nanocrystalline in the nature on the basis of their XRD pattern. The different intense diffraction peak appears at  $2\Theta$  value of  $37.86^{\circ}$ ,  $43.13^{\circ}$ ,  $64.87^{\circ}$ ,  $74.86^{\circ}$ , and  $78.95^{\circ}$ . These  $2\Theta$  values are assigned to the (111), (200), (220), (311), and (222) cubic plane of the developed MgO NPs [\[26](#page-8-24)]. The average crystalline size of the biologically developed MgO NPs is 22.5, which is evaluated by Debye Scherrer's formula (Fig. [2](#page-3-1)).





<span id="page-3-0"></span>**Fig. 1** UV visible absorption spectra of the prepared MgO NPs



<span id="page-3-1"></span>**Fig. 2** XRD pattern of the developed MgO NPs

# **3.3 FTIR**

The bonding and the chemical behavior of the fabricated nanoparticles were evaluated by using FTIR analysis. The FTIR analysis is shown in the fgure. The peaks found at 3425 cm−1 ascribed to the presence of the OH group which is probably due to the presence of the phenolic group (Fig. [3](#page-3-2)). The peak appears at  $2410 \text{ cm}^{-1}$  indicating the presence of the alkynes. The band observed at 1640 cm−1 is probably due to the  $C = C$  stretching of alkenes. A band that appears at  $1380 \text{ cm}^{-1}$  is probably due to the aromatic rings. Moreover, the band at  $2840 \text{ cm}^{-1}$  is assigned to the stretching vibration of the C–H bond of the  $C = C$ . The band that appears at  $863 \text{ cm}^{-1}$  is due to the sulfur compound. The band appears at 685 cm−1 associated with the stretching vibration





<span id="page-3-2"></span>**Fig. 3** FTIR spectrum of the fabricated MgO NPs

of the Mg–O–Mg. Further, the peak that appears at 493 cm<sup>-1</sup> could be attributed to the stretching vibration of the C–Cl bond of the alkyne group [\[27](#page-8-25)[–29\]](#page-8-26).

## **3.4 SEM and EDX studies**

The detailed morphological analysis of the biogenic developed MgO NPs was carried out by using SEM analysis and it indicates that the particles are round with partially agglomerated (Fig. [4](#page-4-0)). The agglomerations of the fabricated nanoparticles are basically due to the van der Waal forces. The developed nanoparticles have diferent particle sizes and the size was found closer to 24 nm. The SEM analysis pictures of the fabricated nanoparticles are shown in the fgure. The surface morphological studies of the fabricated nanoparticles are helpful in the identifcation of their photocatalytic and medical applications [\[30](#page-8-27)].

The fabrication of MgO NPs was further confirmed by EDX analysis. EDX profile of the developed MgO NPs shows strong absorption peak for Mg and O. The other peaks for different elements are most probably present to the phytoconstituents used in the fabrication of MgO NPs (Fig. [5](#page-4-1)).

#### **3.5 The antibacterial study**

The dose-dependent manner is utilized to evaluate the antibacterial efficacy of the developed nanoparticles against the selected bacterial pathogen. The antibacterial potential of newly synthesized MgO NPs is demonstrated in Table [1](#page-4-2).

The microwave-assisted green synthesized MgO NPs signifcantly lethal to the selected microbial organism. The newly synthesized MgO NPs show great antibacterial



<span id="page-4-0"></span>**Fig. 4** The SEM micrograph for the developed MgO NPs



<span id="page-4-1"></span>**Fig. 5** EDX spectrum for the developed MgO NPs

<span id="page-4-2"></span>**Table 1** Antibacterial efficiency of the developed MgO NPs

Name of microorganism	The zone of inhibition measured in mm against different concentration			
		Control $20 \text{ mg/mL}$ $40 \text{ mg/mL}$ $60 \text{ mg/mL}$		
E. coli	25	10	13	14
Bacillus spp.	27	10	14	23
<i>Pseudomonas</i> spp.	26	12	15	16
Klebsiella pneumonia	24	14	16	20
Staphylococcus aureus	23	13	15	21

activity against *Bacillus* spp., *Klebsiella pneumonia*, and *Staphylococcus aureus* and further show less activity against *E. coli* and *Pseudomonas* spp. The result of the antibacterial study also established that the antibacterial potential against all the selected pathogens increased as the concentration of the synthesized nanoparticles was increased (Fig. [6](#page-5-0)).

# **3.6 Mechanistic study of the antibacterial activity**

The synthesized nanoparticles show antibacterial behavior in three well-defned ways: (i) the synthesized nanoparticles are lethal against the bacterial strain because of their small size they readily rapture the cell wall of the bacterial strain, (ii) the entered nanoparticles damage the intercellular structure like mitochondrial vacuoles and ribosomes of the bacterial strain, and (iii) another way which was generally involved in the antibacterial phenomenon of the developed nanoparticles was the production of reactive oxygen species; these species were responsible for cellular toxicity and oxidative stress [[31\]](#page-9-0). This ultimately causes the death of the bacterial strain (Fig. [7\)](#page-6-0).

#### **3.7 The photocatalytic activity**

The photocatalytic efficacy of the developed MgO NPs was attributed to exploring their ability to remove organic dyes like phenol red and rhodamine B from the wastewater released



<span id="page-5-0"></span>**Fig. 6** Disk difusion assay for evaluating the antibacterial potential of the developed MgO NPs against (**a**) *E. coli*, **b** *Bacillus* spp., **c** *Pseudomonas* spp., **d** *Klebsiella pneumonia*, and (**e**) *Staphylococcus aureus*





<span id="page-5-1"></span> $(e)$ 

from various industries. The maximum absorbance of the phenol red and rhodamine B was measured as a time interval. The measurement was carried out at its maximum absorbance. The degradation efficiency of the developed nanoparticles against the phenol red and rhodamine in the presence of solar light and UV light (310 nm) was demonstrated in Figs. [8](#page-6-1) and [9](#page-6-2). The result of the photocatalytic activity indicates that the maximum absorption band of both dyes decreases subsequently with time. According to Lambert–Beer law, concentration is directly proportional to absorbance; hence, the degradation potential can be evaluated by the following equation (Eq. [2\)](#page-5-1).



$$
R = \left\{ (C_0 - C) / C_0 \right\} \times 100
$$
 (2)

where  $C_0$  and  $C$  represent the concentration of dye at time  $t=0$  and  $t=t$ .

# **3.8 Mechanistic study of the photocatalytic behavior of the developed MgO NPs**

The mechanism behind the photocatalytic degradation of dye in solar light and in UV light in the presence of green synthesized MgO NPs is almost similar. The main reaction involved

<span id="page-6-0"></span>

 $C_{\alpha}$ 

 $\mathbf 0$ 

20

40



<span id="page-6-1"></span>**Fig. 8** Photocatalytic degradation of phenol red in presence of MgO NPs

80

Irradiation time(min)

100

120

140

160

60



<span id="page-6-2"></span>**Fig. 9** Photocatalytic degradation of rhodamine B in presence of MgO NPs

<span id="page-6-3"></span>in the photodegradation of dye is the photooxidation of dye by the reactive oxygen species. These reactive oxygen species are generated by electron transfer. When the green synthesized MgO NPs irradiated with light (UV or solar light) electron–hole pair is generated in the MgO nanostructure (Fig. [10\)](#page-6-3). The electron interacts with the molecular oxygen and gives a reactive oxygen species superoxide radical anion and the generated hole interacts with a water molecule to produce hydroxyl free radical species

[\[32\]](#page-9-1). These reactive species are responsible for the photodegradation potential of the developed MgO NPs (Fig. [11](#page-7-0)).

#### **3.9 Antioxidant activity**

Figure [11](#page-7-0) demonstrates the antioxidant potential of the developed nanoparticles. The oxygen donating substance present on the surface morphology of the developed MgO NPs is responsible for the antioxidant behavior of the MgO NPs. The color DPPH solution is changed in this protocol which is the primary indication of the antioxidant behavior of the component (Fig. [12\)](#page-7-1). The synthesized nanoparticles show low antioxidant activity as compared to the ascorbic acid that is used as a positive control in this experiment. The microwaveassisted green synthesized MgO NPs show DPPH radical scavenging activity in the range of 31.41 to 80.13% [\[33](#page-9-2)].



**DNA** Damage

Due to small size nanoparticles

<span id="page-7-0"></span>





<span id="page-7-1"></span>**Fig. 12** Antioxidant potential of the developed MgO NPs

# **4 Conclusion**

In summary, the present research highlights a very efficient eco-friendly low-cost method to develop MgO NPs. The structural characterization of the developed nanoparticles was monitored by standard analytical tools like UV, XRD, FTIR, and SEM analysis. With the help of XRD and SEM analysis, it was stabilized that the particle size of the developed nanoparticles was found to be less than 30 nm and has a spherical shape. The developed MgO NPs have good antibacterial activity against the selected pathogen with the zone of inhibition 23 mm, 21 mm, and 20 mm, respectively, for *Bacillus* spp., *Staphylococcus aureus*, and *Klebsiella pneumonia*. The synthesized nanoparticles also exhibited moderate antibacterial phenomenon against the *E. coli* and *Pseudomonas* spp. The catalytic potential of the synthesized nanoparticles was examined by the degradation of the two carcinogenic dyes phenol red and rhodamine B. The microwave-assisted green synthesized MgO nanoparticles show great antioxidant potential in the DPPH assay. The fnding of our study established that the synthesized nanoparticles have signifcant applications in pharmaceutical industries.

**Author contribution** Waseem Ahmad: conception and design, material preparation, data collection, and analysis. Sanjay Kumar: resources.

**Data availability** The data made available on reasonable request.



#### **Declarations**

**Ethical approval** Not applicable.

**Competing interests** The authors declare no competing interests.

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