

# A Systematic Review of Synthesis MgO Nanoparticles and Their Applications

Hadia Hemmami<sup>1,2,3</sup>, Ilham Ben Amor<sup>1,2\*</sup>, Soumeia Zeghoud<sup>1,2</sup>, Salah Eddine Laouini<sup>1,4</sup>, Emmanuel Chile Nleonu<sup>5</sup>, Pawel Pohl<sup>6</sup>, Jesus Simal-Gandara<sup>7</sup>

<sup>1</sup>Department of Process Engineering and Petrochemical, Faculty of Technology, University of El Oued, El Oued 39000, Algeria.

<sup>2</sup>Renewable Energy Development unit in Arid Zones (UDERZA), University of El Oued, El Oued39000, Algeria.

<sup>3</sup>Laboratory of Applied Chemistry and Environment, Faculty of Exact Sciences, University of El Oued, P.O. Box 789, El Oued 39000, Algeria.

<sup>4</sup>Laboratory of Biotechnology Biomaterials and Condensed Materials, faculte de la technologie, , University of El Oued, P.O. Box 789, El Oued 39000, Algeria.

<sup>5</sup>Department of Chemistry, Federal Polytechnic Nekede, P.M.B. 1036, Owerri, Imo State, Nigeria. <sup>6</sup>Department of Analytical Chemistry and Chemical Metallurgy, Faculty of Chemistry, University of Science and Technology, Wyspianskiego 27, 50-370 Wroclaw, Poland.

<sup>7</sup>Universidade de Vigo, Nutrition and Bromatology Group, Analytical Chemistry and Food Science Department, Faculty of Science, E32004 Ourense, Spain.

**Abstract**: Recently, nanoscale biotechnology has emerged as an essential field of contemporary science and a new era in the study of materials. It draws the attention of many scientists from all over the world due to its versatility in various fields. Many physical, chemical, and biological processes are used to create biomaterials. Among the materials of interest is magnesium oxide (MgO), which can be widely used in medical and biotechnological applications due to its non-toxicity and environmental friendliness. This review article discusses various methods for the synthesis of magnesium oxide nanoparticles (MgONPs), with particular emphasis on recent developments and applications of these nanomaterials.

Keywords: MgONPs, Synthesis techniques, applications.

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## \*Corresponding author's E-mail: <a href="mailto:illambenamor97@gmail.com">illambenamor97@gmail.com</a>

## 1. INTRODUCTION

Nanotechnology is a main driver behind the advancement of different disciplines of study that offers promise for numerous advances. Science and technology, deal with the synthesis, characterization, and development of applications of materials with at least one nanometer-scale dimension (1, 2).

Nanoparticles (NPs) are subgroups of particles with a size range of 1-100 nanometers that make up nanomaterials, which are structural components with a size range of 1-1000 nm (3, 4). Conical, spiral, flat, hollow, and other shapes and structures are just a few examples of the many different forms that can exist. Certainly, they exhibit exceptional physical

traits when compared to their bulk form, which gives them unique mechanical strength, improved stability, and many other advantages, and opens up the possibility for a range of unique applications (5, 6).

MgONPs are gaining more attention than other metal oxide nanoparticles that are frequently used in a variety of fields due to their high strength-to-weight ratio, low density, good functionality, nontoxicity, and hygroscopic properties. Because of their biocompatibility, they are very promising structural materials for implants and other biological systems. These characteristics of MgONPs boost their usefulness in several ways and provide additional benefits. It should not be surprising that MgONPs

have numerous commercial and biological applications in addition to their use in bone regeneration-assisting cryoinjury, antibacterial and antimicrobial inhibition, catalysis, lithium-ion battery production, and elimination of hazardous wastes (3, 7-11).

Because the synthesis method and its course determine the properties of the obtained nanomaterial and its subsequent use, the goal of this review study is to look into the methods involved in the synthesis of MgO NPs, with a particular emphasis on the most recent advancements in their various applications.

# 2. MAGNESIUM OXIDE NANOPARTICLE SYNTHESIS

Several methods are available for producing MgONPs, as shown in Figure 1. The most popular biological and chemical synthesis methods follow the bottom-up approach (12), and they are shown in Figure 1. Specifically, these methods include solve/hydrothermal (13),sol-gel (14), co-precipitation (15), and combustion (16) processes. A special place among these methods is occupied using chemical reactions mediated by the use of plant extracts and microorganism media and fluids, the so-called green synthesis processes (17). In general, the bottom-up approach is advantageous because of its simplicity and ability to control the size and shape of the nanoparticles (18).



Figure 1: Different processes for creating nanoparticles.

# 2.1. Sol-Gel Technique

One of the simplest ways for synthesizing new material structures in the presence of an organic solvent and an inorganic precursor is the sol-gel method (19). Typically, this method is used to produce inorganic compounds like metal oxides and others that are comparable. In the middle of the 19th century, silica gel was first made using the sol-gel process (20). Metal alkoxides can be used to prepare homogeneous solutions, colloidal suspensions (sol), and integrated networks (gel), which, depending on the drying method, can subsequently be converted into xerogels or aerogels.

Mustuli *et al.* (21) focused on the production of nanostructured MgONPs using the sol-gel technique, They discovered that using Mg acetate in combination with a complexing agent in the form of oxalic acid and tartaric acid, the crystal growth could be inhibited.

Sutapa *et al.* (22) also produced MgONPs using Mg acetate. These researchers achieved the creation of cubic-shaped crystals, which were verified, using a scanning electron microscope (SEM), to have the highest texture coefficient value (0.98 in the crystal

plane (222)). They also characterized stress, strain, and crystal energy.

Wahab *et al.* (23), synthesized MgONPs using  $Mg(NO_3)_2$  and NaOH. The sol-gel method described in their work resulted in the production of cubic-shaped MgONPs with a size of 50-60 nm.

In contrast, using Mg ribbons as a precursor, Boddu *et al.* (24) documented the synthesis of MgONPs having a coralline structure. In this case, hydrolysis, supercritical drying, and heat activation processes were carried out after a solution of Mg methoxide was obtained. This applied method resulted in the fabrication of 200–300 nm-sized particles having the aforementioned structure. To produce a nanopowder from MgO xerogel, Mg methoxide was employed by Dercz *et al.* (25) as a precursor, and the hydrolysis in the presence of toluene, followed by the addition of methanol was carried out. In this way, a specific surface area of 138 m<sup>2</sup>/g and an average crystallite size of 7.5 nm were attained for these MgONPs.

Rani *et al.* (26) employed  $Mg(NO_3)_2$ , which dissolved in distilled water. SEM examinations showed that the final particles, produced *via* gel grinding and subsequent annealing, had an average size of 60 nm. Nassar *et al.* (27) employed a mixed sol-gel combustion technique to produce nanostructured MgONPs as well. The scientists discovered that the kind of fuel had no discernible impact on the size and shape of the crystallites when utilizing  $Mg(NO_3)_2$  in combination with oxalic acid, urea, and citric acid (citric acid was used to create the lowest crystallite size, which was about 12 nm).

## 2.2. Co-Precipitation

The synthesis of NPs with this method is based on the idea of precipitation, and frequently uses liquidphase synthesis (28), while vapor-phase synthesis is applied less frequently (29). The precipitating agent is frequently NaOH (30, 31), and two processes, i.e., nucleation and nuclei growth, make up together the homogeneity of the precipitation reaction and the quality of the final nanomaterial product (32). Three basic concepts are taken into account: (i) the diffusion-based single nucleation and the homogeneous growth; (ii) the smaller subunits formation, development, and assemblage; and (iii) the numerous nucleation and the Ostwald ripening growth (33).

Mashad *et al.* (34) produced MgONPs by coprecipitation and evaluated the effects of various reaction parameters, such as temperature, pH, and the molar ratio of precursor (magnesium nitrate), on the quality of the products. They obtained nanoparticles and nanorods with a reasonably high specific surface area (231 m<sup>2</sup>/g for nanoparticles and 176 m<sup>2</sup>/g for nanorods) and a particle size of 50 nm.

Mg(NO<sub>3</sub>)<sub>2</sub> was used as the precursor, while an NH<sub>4</sub>OH solution served as the precipitating agent in work by Kumar *et al.* (35). As a result, MgONPs having an average size of around 11 nm were produced. The impact of the polyethylene glycol (PEG) content on the characteristics of MgONPs produced by the coprecipitation method was also investigated by Karthikeyan *et al.* (36). In this work, Mg(NO<sub>3</sub>)<sub>2</sub> was used as a precursor while NaOH as a precipitation agent. When PEG was additionally used, it resulted in doubling the size of crystallites, according to XRD measurements (8.6 nm *vs.*14.8–15.9 nm without and with PEG, respectively). What is more, PEG-modified MgONPs were spherical.

By calcining MgCO<sub>3</sub>, which Frantina *et al.* (37) initially obtained by combining  $(NH_4)_2CO_3$  and MgCl<sub>2</sub>, MgONPs were also produced. The XRD data suggested a cubic structure with an average crystallite size of 24 nm. With negligible changes in the particle size, the spherical shape of the resulting MgONPs was confirmed using SEM, while their average size was 50.9 nm.

MgONPs reported by Kushwaha *et al.* (38) were produced by several distinct techniques (sol-gel, solution combustion, and a solution of cetyltrimethylammonium bromide (CTAB)), as well as the co-precipitation technique. Their findings demonstrated that 4.9 eV bandgap MgONPs were produced using the chemical method. The crystallite size was shown by XRD to be 14.8 nm, and the authors also reported a hydrodynamic particle size of 100 nm. MgO nanotubes were also prepared by Tandon and Chauhan (32) using Mg acetate and NaOH. Using XRD, the average crystal size was calculated to be 34 nm. The field emission (FE)SEM data showed that the resulting nanomaterial had a tubular shape, with an inner diameter of 31 nm and an estimated outside diameter of 78 nm. Additionally, a greater bandgap of MgONPS in comparison to the prior example (5.73 eV) was reported.

## 2.3. Combustion Technique

Due to its effectiveness and affordability, the combustion process is commonly utilized to manufacture metal oxide nanoparticles (39). Two strategies can be used for that, including so-called "volume "self-propagating synthesis" and the combustion synthesis" (40). In the case of "selfpropagating synthesis", the production of solid products occurs without the need for the energy input (41), because spontaneous redox reactions occur between the precursor (oxidizer) and the reductant (fuel) combined at the molecular level in solution. These reactions are initiated by an outside source. In the second case ("volume combustion synthesis"), the sample is heated until the reaction starts, spreading across its whole volume. The latter approach is especially recommended for moderate exothermic reactions that require preheating before the ignition, even though this sort of preparation is more difficult to control (42).

Accordingly, when urea was used as a fuel and  $Mq(NO_3)_2$  as an oxidizer, it was possible to produce MgONPs with a cubic structure and a crystallite size of around 22 nm, according to the XRD data(43). The resulting MgONPs were uniformly sized and spherical, as established by SEM. Interestingly, the synthesized nanomaterial exhibited a bandgap of just 2.9 eV, which is much lower as compared to earlier research. The same raw ingredients were utilized by Rao et al. (44), who tested the impact of the fuel-tooxidizer ratio on the quality of the produced nanomaterial. The findings showed that except for the percentage of oxidizer was 0.75, manufactured MgONPs have larger crystallite sizes (18–53 nm) when the fuel contribution increases. Changes in the burn rate, enthalpy, or ignition temperature might be responsible for this.

Ranjan et *al.* (45) reported a variation of this procedure, using  $Mg(NO_3)_2$  as the precursor and glycine as the fuel. The estimated crystallite size, based on the XRD measurements, was 20.8 nm in this case.

On the other hand, Therami *et al.* (46) used citric acid as fuel. The authors examined how the specified parameters of MgONPs were impacted by the concentration of this acid. The most significant changes obtained were observed for the particle size of MgONPs (it decreased from 35 to 20 nm), their bandgap (it increased from 4.72 to 5.35 eV), and their morphology (vacuolar, flower-like, and flake-like).

MgONPs were also obtained by Kumar *et al.* (47), who used  $Mg(NO_3)_2$  and a parthenium plant extract. The primary objective was to analyze the impact of the fuel quantity on the bandgap width (5.3-5.45 eV) and the crystallite size (27-35 nm). However, the variation of both parameters was not as pronounced as it was reported in other works.

#### 2.4. Solvo- and Hydrothermal Method

Another popular technique for regulating the formation of crystals in a variety of materials is the solvothermal approach (48). The required products are generated when a precursor and a suitable solvent are put in an autoclave and exposed simultaneously to a high temperature and a high pressure (49). In contrast to the co-precipitation approach, these reaction conditions (temperature and pressure) enable to obtain the high crystallinity materials (50). Solvents other than water are typically utilized in the "solvothermal" technique, including alcohols or other organic or inorganic solvents. When water is used as the solvent, this process can be referred to as "hydrothermal."

Devaraja *et al.* (51) used Mg(NO<sub>3</sub>)<sub>2</sub> and NaOH to produce a nanocrystalline MgO powder and evaluated its quality. The MgONPs produced by them were porous, with an average crystallite size of 25 nm and an optical energy bandgap of 5.5 eV. Al-Hazmi *et al.* (52), on the other hand, obtained nanofibers by the direct interaction of urea and Mg acetate. These fibers had an average crystallite size of 6 nm, which corresponded to their diameter, and the length of 10 nm, as measured by TEM.

Ding et al. reported the synthesis of rod- and tubeshaped  $Mq(OH)_2$ , which was subjected to thermal decomposition to fabricate MgONPs (53). The author's findings demonstrated that the hydrothermal approach could be used to manipulate the crystallite size of the resultant MgONPs, and their shape and structure. The material for the synthesis was either Mg powder,  $Mg(SO_4)_2$ , or  $Mg(NO_3)_2$ . Due to various experimental circumstances, numerous morphologies (lamellar, needle-like, and rod-like) of MgONPs were achieved. The resultant nanomaterials were 20 to 600 nm in size and had a specific surface area of more than 100  $m^2/g$ .

Rukh *et al.* (54) used Mg powder as the precursor for the MgONPs synthesis. The reaction medium for the synthesis was a mixture of  $H_2O_2$  and de-ionized water. Using this method, MgONPs with an 18 nm crystallite size were obtained. Nanoplates are another form of nanostructured MgO that was reported to be produced by Duong *et al.* (54). Additionally, the scientists utilized sodium dodecyl sulfate (SDS), PEG, CTAB, and Mg(NO<sub>3</sub>)<sub>2</sub> to regulate the morphology of the resulting nanomaterials. MgONPs produced using the hydrothermal process in conjunction with SDS were the most intriguing since they had the largest specific surface area (126 m<sup>2</sup>/g) and the ideal disc shape (thickness 5 nm, diameter 40–60 nm).

#### 2.5. Green Synthesis of MgO Nanoparticles

Researchers have shown a growing interest in the production of MgONPs through biological processes over the past ten years. The development and significance of this synthesis type are mostly related to the possibility of the use of much fewer chemicals, making this less cost-effective and more environmentally friendly (55-57).

The traditional chemically- or physically-based techniques for the synthesis of MgONPs are less practical and less ecologically benign than their biologically-based alternatives, also known as (56, 58). Consequently, the term "green synthesis" is frequently used to describe biological techniques. The large-scale synthesis of nanoparticles utilizing the green methods is always a difficult undertaking, and these are only performed at laboratory-scale processes. However, thanks to advancements in the understanding of the nature of the biological extract composition and the reaction with metal ions, large-scale preparation may soon be feasible without the need for any powerful machines (59).

To lessen the hazardous nature of the nanoproducts, biological substrates such as plants, bacteria, algae, and fungi are frequently utilized in place of chemical compounds used as stabilizers and solvents (59). The greener way to produce MgONPs involved the use of the available precursors, such as  $Mg(NO_3)_2$ , MgCl<sub>2</sub>, Mg acetate, and Mg(SO<sub>4</sub>)<sub>2</sub>, and different biological agents, including plants, microorganisms, and biomolecules. The precursor was combined with the previously produced biological extracts of plants, microorganisms, or templates to prepare homogeneous mixtures, which were then subjected to thermal treatment (60-62).

There are several papers on the production of MgONPs. The information on the various synthetic processes used to produce nanostructured MgO reported in the literature is given in Table 2. Many biological templates were employed for the production of MgONPs, as can be seen from the data given. The production of MgONPs with various sizes and morphologies was ultimately achieved by the variation in the reaction time, the concentration of the Mg precursors, pH, and the temperature of the reactants.

For instance, it was established that the particle size of MgONPs grows along with the increasing dose of the biological substrate(63). According to the Mie hypothesis, the absorbance of light is proportional to the particle size of the metal nanoparticles formed. However, it was discovered (64) that when the extract of Amaranthus tricolor and Mg acetates  $(Mg(C_2H_3O_2)_2)$  were used, the MgONPs produced were found to deviate from the Mie theory, showing the lower absorbance and the increased particle size when the amount of the biological substrate was higher. When they added 5 mL of the Amaranthus tricolor extract and Mg acetate, they found that the product had a little variation from the Mie theory; it had decreased absorbance and increased particle size. Smaller-sized MgONPs resulted from the addition of 10 and 15 mL of the Amaranthus tricolor

extract to the same precursor. This demonstrated how the presence of the biological substrates affects the size of MgONPs. According to scientists, the addition of capping agents could also improve the production of MgONPs. Smaller amounts of the capping agents caused the particles to adhere to one another and, to larger extents, the formation of larger particles. It is also obvious that the use of lower concentrations of the biological substrates would produce more homogeneous and bigger nanoparticles, whereas higher concentrations of the biological substrates would produce less stable nanoparticles. It is also evident that the presence of flavanoids in the biological substrates is responsible for these modifications. The development of MgO NPs is also influenced by reaction time. Additionally, the size of the product rises along with the response time. Numerous pH reports at 3.2, 5.2, 7.2, and 9.2 are available. It is clear from examining the impact of pH reports that a pH of 3.2 results in good particle size. The biological templates' active components, like flavonoids and polyphenols, are particularly effective at reducing metal ions at this pH level (65).

Khan et al. (66) noticed that the temperature also has an impact on the formation of MgONPs. The authors conducted the reaction at temperatures between 30 and 70 °C while holding all other variables constant. Generally speaking, the appropriate temperature should be kept for the formation of MgONPs with phytochemical support (Flavonoids, phenols, and other species). Otherwise, the phytochemicals utilized to synthesize MgONPs could alter their structure. The highest absorbance (81%) range of methylene blue (MB) dye was achieved at 30 °C. Additionally, it was noticed that when the temperature was raised, product agglomeration occurred, while product formation was prevented. Furthermore, the production of the final product was hampered by precursor concentrations greater than 0.001 mol/L. The final loss of chemicals, energy, and time will occur from the addition of an excessive concentration of the precursors. The physicochemical properties of MgONPs can be altered depending on the conditions of the green synthesis conditions (see details in Table 2).

<b>Table 1:</b> Individual ingredients and reaction circumstances for the bottom-up production of nanostructural							
MgO using various techniques.							

Sol-Gel Technique									
Precursor	Solvent(s)	Drying temperature (°C)	Temperature of calcination (°C)	Time of calcination (h)	Size (nm)	Expected applications	Ref.		
Mg(NO <sub>3</sub> ) <sub>2</sub>	C <sub>6</sub> H <sub>8</sub> O <sub>7</sub> , C <sub>2</sub> H <sub>2</sub> O <sub>4</sub> , NH <sub>2</sub> CONH <sub>2</sub>	350	550, 800	2	12	catalyst	(27)		
Mg(CH <sub>3</sub> COO) <sub>2</sub>	C <sub>4</sub> H <sub>6</sub> O <sub>6</sub> , NaOH	-	600	6	-	-	(21)		
Mg(CH <sub>3</sub> COO) <sub>2</sub>	$C_2H_2O_4$	200	950	6	-	-	(22)		
Mg(NO <sub>3</sub> ) <sub>2</sub>	NaOH	300	500	2	50-60	adsorber	(23)		
Mg(OCH <sub>3</sub> ) <sub>2</sub>	CH <sub>3</sub> OH, C <sub>7</sub> H <sub>8</sub>	-	500	5	200-300	-	(24)		
Mg(OCH <sub>3</sub> ) <sub>2</sub>	CH <sub>3</sub> OH;C <sub>7</sub> H <sub>8</sub>	60	450	-	Ca.8	-	(25)		
$Mg(NO_3)_2$	H <sub>2</sub> O	150	500	2	60	-	(26)		
	T	Co-F	Precipitation Tech	nique					
Precursor	Precipitation agent	Reaction temperature (°C)	Temperature of calcination (°C)	Time of calcination (h)	Size (nm)	Expected applications	Ref.		
Mg(NO <sub>3</sub> ) <sub>2</sub>	NaOH	Room	Room	-	78	antibacterial agent	(32)		
Mg(NO <sub>3</sub> ) <sub>2</sub>	NaOH	-	440	4.5	-	catalyst	(38)		
$Mg(NO_3)_2$	NH₄OH	60, 70, 80	550	2	50	-	(34)		
Mg(NO <sub>3</sub> ) <sub>2</sub>	NH₄OH	100	600	4-6	11	antibacterial agent	(35)		
Mg(NO <sub>3</sub> ) <sub>2</sub>	NaOH	Room	500	4	14-16	antibacterial agent	(36)		
		Co	ombustion Techni	que					
Oxidizer	Fuel	Ignition temperature	Temperature of calcination (°C)	Time of calcination (h)	Size (nm)	Expected applications	Ref.		
	Parthenium	400	-	-	27-35	photocatalyst	(47)		
	extract	70-80	500	3	22	adsorber	(43)		
$M_{II}(NO_{2})_{2}$	NH <sub>2</sub> CONH <sub>2</sub>	100	300	2	18-53	-	(44)		
119(1103)2	NH <sub>2</sub> CH <sub>2</sub> COOH	170	600	2	Ca.21	fuel additive	(45)		
	C <sub>6</sub> H <sub>8</sub> O <sub>7</sub>	100	400	15 min	20-35	antibacterial agent	(46)		
	1	Solvo- a	nd Hydrothermal	Technique	r				
Oxidizer	Solvent	Autoclave temperature	Temperature of calcination (°C)	Time of calcination (h)	Size (nm)	Expected applications	Ref.		
$Mg(NO_3)_2$	NaOH	100	500	4	40-60	adsorber	(67)		
Mg	H <sub>2</sub> O <sub>2</sub>	220	-	-	18	antibacterial	(54)		
MgSO <sub>4</sub>	NH <sub>3</sub> H <sub>2</sub> O;en- H <sub>2</sub> O	180	280-450	1;2	100-200	catalyst	(53)		

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Mg(NO <sub>3</sub> ) <sub>2</sub>	NaOH	130	400-800	2	25	-	(51)
Mg(CH₃CÓO)	NH <sub>2</sub> CONH <sub>2</sub>	180	600	1	6	antibacterial	(52)
$Mg(NO_3)_2$	NaOH	80 280-450 1;2 50		50	catalyst	(53)	
Green Synthesis							
Precursor solution	Precursor Reagent solution extract		Temperature of calcination (°C)	Time of calcination (h)	Size (nm)	Expected applications	Ref.
	Nephelium Iappaceum	80	450	-	55	-	(68)
Mg(NO₃)₂	Trigonella foenum- graecum	80	600	4	14	antibacterial agent	(69)
	Rosa foribunda	90	-	-	10	antibacterial agent	(70)
Bulk MgO	Rosmarinus offcinalis	70	-	-	8.8	antibacterial agent	(71)
	Dalbergia sissoo	30-70	-	-	50	photocatalyst	(66)
Mg(NO₃)₂	Saussurea costus	80	450	3	30	photocatalyst	(72)
	Swertia chirayaita	55	400	4	< 20	antibacterial agent	(73)
MgCl <sub>2</sub>	Moringa oleifera	90	600	5	21	antibacterial agent	(74)
Mg(NO <sub>3</sub> ) <sub>2</sub>	Tecoma stans	90	550	6	20-50	adsorber	(75)

**Table 2:** Reported processes for making MgO NPs through green synthesis.

Material used	Particle size (nm)	Morphology of nanomaterial	Activity carried	Ref.
<i>Citrus limon</i> leaf extract	12-80	nanoflakes	nil	(58)
Rosmarinus officinalis	<20	nanoflowers	antibacterial activity	(71)
Nephelium lappaceum	60-70	cubic	nil	(76)
Solanum trilobatum	30 and 42	spherical	antibacterial and antioxidant activity	(77)
Mucuna pruiens seeds	35	spherical	antibacterial and photocatalytic activity	(78)
Rhizophora lamarckii	20 and 50	hexagonal and spherical	antibacterial activity	(79)
Aloe vera	8.6	dense rock-shaped flakes	antibacterial and photocatalytic activity	(80)
Amaranthus blitum and aloe vera	26-50	spherical	water treatment	(81)
Mushroom extract	20-15	cubic	seed germination	(82)
Aspergillus tubingensis	2.8	sphere	nil	(83)
Aspergillus niger	43-91	sphere	antibacterial activity	(84)
Lactobacillus plantarum Lactobacillus sporogenes	30	cubic	anticancer activity	(85)
Aspergillus fumigatus	0.3-94	nil	nil	(86)
Manihot esculenta	37	hexagonal	nil	(87)
Sargasssum wightii	69	cubic	antimicrobial and photocatalytic activity	(88)
<i>Artemisia abrotanum</i> herb	10	clusters	photocatalytic and antioxidant activity	(89)
Rhododendron arboretum	nil	nil	antibacterial activity	(62)
Ocimun sanctum	50-100	nanoflakes	antibacterial and antioxidant activity	(81)
Chamaemelum nobile flower extract	20-40	nanoflakes	insect repellent	(90)
Curcumin	35	rod-like and spherical - like shape	catalytic properties	(91)
Acacia gum	50-78	nanoflowers, cubic	catalytic properties	(92)
Brassica olercea and Punica granatum	30-65	spherical	photocatalytic and anticancer activity	(93)

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Orange peel extract	>10	spherical	antibacterial and antimicrobial activity	(94)
Curry leaves	20	spherical	photocatalytic activity	(95)
Swertia chirayaita	<20	spherical	antibacterial activity	(73)
<i>Pisidium guvajava</i> and <i>aloe vera</i>	50-90	cubic	antibacterial activity	(96)
Lepidium sativum	33	nanoflakes	photocatalytic activity	(97)
Matricaria chamomilla L. extract	18 and 16	disc-shape	antibacterial activity	(98)

#### 3. RESOURCES THAT CAN BE USED AS A MAGNESIUM SOURCE TO PREPARE MgO NPs

MgO nanoparticles with superior properties are important in industry, and these particles can be prepared from different sources using different methods. Table 3 shows the resources that can be used to manufacture MgO NPs as a source of magnesium in a direct way. As for the indirect method, an intermediate is extracted and subsequently treated to produce magnesium oxide(Table 4). The following chemical equations (1-4) explain the mechanism of transformation of the medium into magnesium oxides.

**Table 3:**There are several resources that may be employed to manufacture MgO NPs as a magnesium source.

Resources	Presses	Intermediate reactant	Temperature of calcination (°C)	Yield%	Purity%	Size (nm)	Ref.
Dolomite CaMg(CO <sub>3</sub> ) <sub>2</sub>	Pyrohydrolysis process	HCI CO <sub>2</sub>	600	98.10	98.86	100	(99)
Dolomite: CaMg(CO <sub>3</sub> ) <sub>2</sub> ]	Pyrohydrolysis process	HCI	FOO		95.48		(100)
Dolime (CaO.MgO)	Pyrohydrolysis process	MgCl <sub>2</sub>	500		87.80		(100)
Serpentinite;		(NH <sub>4</sub> ) <sub>2</sub> SO <sub>4</sub> NaHSO4	1300		high purity	62	(101)
Dolomite : poly(acrylate) magnesium hydroxide	Pyrohydrolysis process	poly(acrylate) Na <sub>2</sub> S <sub>2</sub> O <sub>8</sub> MgCl <sub>2</sub> NaOH	500		high purity	20	(102)
Sea water	Pyrohydrolysis process	MgCl₂ NaOH	1400	43	56		(103)

Table 4: Various resources that may be	e employed to prepare an intermediate reactant using a magnesium
	source.

Resources	Technique	Reactants	Temperature of reaction (°C)	Yield %	Purity %	Size (nm)	Ref.
serpentinite mineral	precipitation	NH₄OH HNO₃	80		high purity	30	(104)
Serpentine 4MgCO3.Mg(OH) <sub>2</sub> . 4H2O	precipitation	NH <sub>3</sub> .H <sub>2</sub> O NH <sub>4</sub> HCO <sub>3</sub>	40~70	96.3	high purity		(105)

 $MgCl_2.6H_2O \xrightarrow{\Delta} MgO_{(s)} + 2HCl_{(g)} + 5H_2O \dots \dots \dots (1)$ 

$$MgCl_2. H_2O_{(s)} \rightleftharpoons MgCl_{2(s)} + H_2O_{(v)} \dots \dots \dots (2)$$

 $MgCl_2$ .  $H_2O_{(s)} \rightleftharpoons MgOHCl_{(s)} + HCl_{(g)} \dots \dots \dots (3)$ 

 $MgOHCl_{(s)} \rightleftharpoons MgO_{(s)} + HCl \dots \dots (4)$ 

# 4. CHARACTERISTICS MgO NPs

Magnesium oxide may also be identified and structurally characterized using Fourier transform infrared spectroscopy (FTIR)(106, 107). Depending on the wavelength of the incoming light, transmittance or absorbance are the most typical interpretations. Figure 2a displays typical FTIR spectra for magnesium oxide in the absorbance mode. Mono-coordinated hydroxyl (-OH) groups are indicated at 3100-3500 cm<sup>-1</sup>. The stretching frequency of H-O-H is linked to a wide band at roughly 3461 cm<sup>-1</sup>. At roughly 673 cm<sup>-1</sup>, the wide band stretching vibration matches the Mg-O stretching vibration. The stretching vibration of magnesium oxide is shown by a prominent peak with a center of 433-769 cm<sup>-1</sup>.

Using X-ray diffraction (XRD) (JCPDS Standard No. 01-089-7746), it is possible to characterize the crystal structure of magnesium oxide. Important peaks can be assigned near the 20 values of 36.8, 42.9, 62.19, 74.6, and 78.58°, which can be indexed to the lattice planes (111), (200), (220), (311), and (222), respectively. Comparable findings have been frequently reported(108) (Figure 2b). The most

popular method in the literature for interpreting XRD data to calculate the crystallite size of produced nanostructures is the Scherrer equation(109). Nevertheless, the average crystallinity size up to around 200 nm(110) is the limiting factor in terms of employing the Scherrer equation. This is caused by the fact that when crystallite size increases, the diffraction peak's widening reduces(111). Hence, it is challenging to distinguish between the widening of the peak caused by crystallite size and the

broadening caused by other causes (e.g., size distributions and shape of the crystallites)(106).

In addition to the methods mentioned above, UV-Vis spectroscopy (UV-Vis) and spectra are taken between 200 nm and 800 nm. They may also be used to analyze MgO, especially to establish the bandgap energy(112, 113). Tauc plots and absorbance spectra are both utilized for this. To measure the bandgap width, the photoluminescence technique(114) is usually employed.



**Figure 2:** FTIR spectra (a) and an XRD pattern(b) used to characterize the structural properties of MgONPs (redrawn and adapted from the results presented in(115)).

Depending on several variables, including the synthesis process, the synthesis circumstances, and the post-treatment techniques, MgO structures can have a variety of morphologies. Typical morphological variations between MgO formations include the following:

**Particle size and shape:** The synthesis technique utilized can have a significant impact on the size and form of MgO particles. For instance, sol-gel processes may yield a range of morphologies, including nanowires and sheets, whereas precipitation methods usually produce spherical particles with a restricted size distribution(116).

**Surface area:** Depending on their form, MgO structures' surface areas can also change. Larger particles often have lower surface areas than nanoparticles or nanowires, which might be crucial for applications like catalysis(117).

**Porosity:** Depending on the synthesis technique employed, MgO structures can be either very porous

or non-porous. For instance, although hydrothermal synthesis may yield structures with a high degree of crystallinity and little porosity, flame synthesis can produce very porous structures, such as hollow spheres(118).

**Crystal structure:** The crystal formations of MgO can be cubic, hexagonal, or tetragonal. The material's chemical and physical characteristics can be affected by the crystal structure(119).

**Surface chemistry:** Depending on their form, MgO structures' surfaces can have different chemical compositions. For instance, exposed crystal planes or surface flaws might influence the reactivity of MgO in catalytic processes(120).

The morphology and properties of the prepared MgONPs differ and depend on the synthesis route and processing conditions. Figure 3 illustrates the various morphologies that can be seen in MgO nanoparticles.



**Figure 3.** Different morphologies of MgO NPs: (a) flower-like(121), (b) spherical(115),(c) irregular shaped flakes(97),(d) Clusters(122),(e) Hexagonal(87),(f) flakes shape(87).

#### **5. APPLICATIONS**

MgONPs have a large specific area, a broadband gap, and good physical and chemical properties, providing them with strong technological applications. Additionally, MgONPs exhibit a propensity to generate superoxides when they are in contact with  $O_2$  found in bacterial cell walls. The bacterial cell walls and their phospholipids are promptly destroyed by these superoxides, which are highly responsive to them. MgONPs can also be used in many applications, as displayed in Figure 4, including the production and/or modification of fuel cells, ceramics, batteries, supercapacitors, and electronics, in addition to environment and agriculture purposes.



Figure 4: Different applications of MgONPs.

#### 5.1. Energy

Recently, the global world has seen a surge in air pollution, global warming, and sea level rise as a result of the depletion of fossil fuels. Therefore, it is critical to locate a substitute for these changes. Fuel cells, solar cells, and batteries as fossil fuel alternatives could be a very good remedy. In addition to the production of  $H_2$ , which is a good fuel and a superior substitute for carbon-based bi-products, they will also release water as a byproduct. Magnesium performs better than other metals in the storage of  $H_2$ . The storage of  $H_2$  is stated to benefit from the use of chemical hydrides such NaAlH<sub>4</sub>, LiBH<sub>4</sub>, and LiNH<sub>2</sub> as well as metal hydrides like Pd@H, V@H, alloys like TiFeH<sub>2</sub>, LaNi<sub>5</sub>H<sub>6</sub>, Ti@V@ Mn@H, Mg<sub>2</sub>NiH<sub>4</sub>, and certain complicated hydrides. Compared to the other hydrides, Mg has the benefit of being extremely abundant in the earth's crust, having a higher capacity to store H<sub>2</sub>, and having an ecologically benign and cost-effective nature (123). There are numerous reports of these MgO-based batteries that use polymers (124), pigments like acetylene black (125), metals like Na (126), Li, V (127), carbon substrates (128), and B. These energy storage technologies are reliable, safe, affordable, and ecologically benign.

# 5.2. Catalysis

MgONPs are also frequently used in heterogeneous catalysis for several chemical reactions, including the oxidative coupling of CH4, the dehydration of the benzylation of aromatics, alcohols. the dehydrohalogenation of halogenated hydrocarbons, the production of pyranopyrazole and its derivatives, the benzylation of aromatics, the dehydrohalogenation of halogenated hydrocarbons. In this case, nano-structured MgO is utilized as a support for catalysts due to its structure, basicity, and electronic and electrochemical properties, making it easier to move the electrons across the catalyst surface (129, 130). Vegetable oils were recently trans-esterified using MgO as a catalyst (131). Other reactions, *i.e.*, Wittig (132), Cyanosilylation (81), Aldol (133), Mannich (134), aza-Michael (135), Baylis-Hillman (136, 137), were also catalyzed with the aid of MgONPs. The enormous surface area and the distinctive morphology of MgO are thought to be the cause of its observed high catalytic activity (138).

# 5.3. Agricultures

MgONPs are also known to offer several benefits, including low phytotoxicity, non-genotoxicity, and non-biotoxicity to people, and thermal stability, which open up a wide range of potential applications of this nanomaterial for plant protection (139). Along with the aforementioned qualities, MgONPs also possess several additional traits that make them particularly useful in a variety of different agricultural applications, as illustrated in Figure 5 (140). Additionally, these nanoparticles aid in the development of seedling and plant growth and are utilized as an authorized food supplement, a food additive, a color retentate, and also in increasing the agricultural production of peanuts (141, 142).



Figure 5: Potential applications for MgO NPs in the agricultural domain.

# 5.4. Biomedical

Because of their high absorption capability, high reaction activity, active catalysis property, and enzyme immobilization, MgONPs are used in the development of the diagnosis of cancer and the guidance of the cure plan through medical imaging. For use in dentistry, surgery, bacterial suppression, tissue engineering, and bone mending, bioactive glass is currently being developed(143, 144). Research findings support the addition of MgONPs to several medicinally valuable chemicals due to their various qualities, including antibacterial, anticancer, biocompatibility, nontoxicity, and cheap cost. MgO also seems to have several safe and useful medicinal applications. Given the potential for negative impacts from exposure to MgONPs, we must have the best method to both reap the benefits of MgONPs and prevent any negative effects that may arise(145).

# 5.5. Anti-microbial Activity

By inducing a breach in their cell membrane and eventually resulting in their death, MgO NPs have antibacterial action against food-borne pathogens such as E. coli and Salmonella entiritidis (146). The bacterial strain Acidovorax oryzae is the source of the illness known as the bacterial brown stripe, which is known to spread among rice and entirely ruin rice farming. As a result, Ogunyemi et al. reported on the biosynthesis of MgO NPs utilizing Matricaria chamomilla L., which showed a good inhibitory impact on the development of Acidovorax oryzae bacteria (98). The worst wilt disease in R. *solanacearum* is caused by phytopathogenic bacteria, which were found to have a favorable antibacterial response to MgO NPs in another study (147). Accordingly, MgO NP nanoflowers were also developed because they possess the ability to inhibit bacterial infections and shield crops from harmful attacks (148). Due to the role of Mg in the

pathogenesis, plant defense, and other physiological processes, as well as the importance of this metal for maintaining balanced nutrition in plants, MgO NPs are known to have a strong antifungal action even at low fungicidal concentrations (149-152).

## 5.6. Anti-biofilm Activity and Anti-insecticidal

Cry genes are proteins produced by Bacillus thuringiensis that function as an insecticide against a variety of insects, including nematodes. However, this protein is released into the soil by water, which hinders its insecticidal effectiveness. By adhering this protein to MgO NPs and subsequently transferring them to the surface of cotton leaves, Rao et al.(153) indicated increased insect fatality rates across the board. As a result, cry protein was transferred via NPs, which enhanced MgO their use as bioinsecticides (154). By limiting the development of the biofilms, MgO NPs are known to promote systemic resistance against the gram-negative, plant-pathogenic bacterium R. solanacearum. They can also start the signaling of pathways for phytohormones like jasmonic acid and salicylic acid, which are crucial components of the plant's defense mechanisms. Therefore, it should be noted that MqONPs are highly efficient anti-microbial, agents, anti-insecticidal antibiofilm and in agricultural fields (155, 156).

# 5.7. Environmental

The major issue of environmental pollution affects both developed and developing countries worldwide. There are several ways to deal with this global problem, but one downside is that certain cleaning chemicals have side effects that make them act as contaminants themselves. Nanoparticles appear to be a fantastic replacement for several various environmental applications (Figure 4). MgONPs are effective in a variety of environmental remediation. These metal oxide NPs are utilized as a possible adsorber of harmful gases, including NO<sub>2</sub> and SO<sub>2</sub>, due to their strong adsorbing characteristics, wide surface area, and high reaction capacity (157-160).

## 5.7.1. Dye removal

2,4-Dichlorophenol (2,4-DCP) is a hazardous substance that is often discharged from paper companies into various water sources, functioning as a main effluent in water. This chlorophenol compound (2,4-DCP) is known to have negative effects on people, animals, and plants who consume it, but treatment with MgO NPs has resulted in its rapid degradation because magnesium oxide acts as a catalyst in the degradation of this dye through the ionization Technique (161).

Acid Red 73 dye, a water contaminant released in large quantities from the textile industry, was removed by S. Jorfi et al., and B.J.H. Ng et al. showed that the activity of ferrate VI, which oxidizes the Blue 203 dye (a water-contaminating dye from the leather and cosmetic industries), is enhanced by MgO NPs (162). Numerous other industries, like the fabric and clothing sectors, employ a variety of dyes to color clothing, with indigo carmine, a water effluent, being one of the most often used ones. MgO NPs were created by A. Bagheri GH et al. and used

as a photocatalyst for the photocatalytic decolorization of indigo carmine(163).

# 5.7.2. Heavy metal ion removal and detection

MgO NPs have been shown by Y. Cai et al. to be a novel possibility for the removal of heavy metals like lead ( $Pb^{2+}$ ) and cadmium ( $Cd^{2+}$ )(164). Nanocomposites such as magnesium oxide-copper oxide nanocomposites and magnesium oxidemanganese oxide nanocomposites, which compete with other nanoparticles involved in the removal of heavy metals from water, are effective adsorbents and have demonstrated high adsorbing properties towards heavy metal ions such as lead, arsenic, and mercury(165).

Improved MgO NPs' sensitivity, By demonstrating the exceptional detection of heavy metals like nickel, copper, and cadmium that are present in significant concentrations in well water, tap water, as well as seawater, these nanoparticles, when modified with graphene oxide, demonstrate (166).

## 5.7.3. Chemical toxin detection and elimination

Magnesium oxide nanoparticles' ability to act as detoxifying agents is widely exploited in many different contexts, one of which is the identification and elimination of chemical pollutants. When treated with MgO NPs, the highly toxic chemical bis(2chloroethyl) sulfide, also known as sulfur mustard and typically used as a biological warfare agent, can be broken down into non-harmful products like divinyl sulfide, thiodiglycol, and 2-chloroethyl vinyl sulfide, which are the byproducts of elimination and nucleophilic substitution reactions, respectively (167). According to S. Ali et al., 2,4,6-trinitrophenyl, a very hazardous pollutant known to induce tumors, liver malfunctions, skin-related problems, etc., is degraded in MgO NPs, and ZnO NPs(168).

# *5.7.4. degradation of pesticides and hydrogen peroxide sensors*

Although pesticides serve to increase agricultural productivity by preventing pests and insects from destroying crops, when these chemicals leak from the field into other water sources, they cause several dangerous illnesses to both plants and people. According to L.E. Lange et al., etching MgO NPs combined with polypropylene improves the chemical stability of the reactive sites present in the nanoparticles that break down methyl parathion, an organophosphate insecticide (169). Aluminum oxide and MgO NPs have been discovered to reduce the harmful effects of the diazinon herbicide, which the Environmental Protection Agency has prohibited due to its high toxicity toward both plants and people(170).

Hydrogen peroxide, a consequence of highly selective oxidative processes, has numerous important uses in a wide range of industries and disciplines, including medicinal, therapeutic, environmental, agricultural, industrial, and many more. As a result, its identification by a sensitive and precise approach is required(171). MgO NPs, an inorganic substance, and chitosan, an organic polymer, are used to create biosensor devices for the

detection of hydrogen peroxide(172). Nanosensors are created for the detection of hydrogen peroxide in milk using magnesium oxide nanoparticles. The created nano biosensor is inexpensive, quick, and very sensitive, able to pick up even the tiniest amount of  $H_2O_2$  (173).

Despite the numerous uses for MgO NPs already mentioned, there are still many more. For example, very small amounts of these particles are sufficient to improve the ability of polyurethane films to resist

and particles, corrosion. these along with nanofiltration membranes, can remove pollutants like nitrogen species, organic matter, bacteria, heavy metals, and suspended solid particles to make water safe for drinking(174). Due to their unique qualities, repeatability, such as high recovery and electrostatic, attraction abrasiveness, and oxidizing power, which work together to increase biocidal capabilities, these nanoparticles have become widely used in wastewater treatment in Figure 6 (175).



Figure 6: Environmental applications for MgO NPs.

## 6. CONCLUSION

Physical, chemical, and biological approaches to the synthesis of MgONPs are surveyed in the present review. Chemical and physical methods typically utilize toxic materials and are related to high energy consumption. Biological methods are currently being advocated by researchers due to their simplicity, cost-effectiveness, and environmental friendliness. Therefore, it is important to put a greater emphasis on more widespread greener methods to produce MgONPs, since this activity will be associated to some extent with the reduction of environmental pollution. MgONPs have various industrial applications, which can change energy production and protect crops from diseases caused by plant pathogens. The major problem of nanostructured MgO, as synthesized using different routes, is the occurrence of a wide band gap. Therefore, there is a crucial need to develop the synthesis method that will enable obtaining MgONPs of a narrow bandgap, making a much wider industrial application of this nanomaterial.

## **7. CONFLICT OF INTEREST**

The authors possessed no relevant financial or non-financial interests.

## 8. ACKNOWLEDGMENTS

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