# Research Article



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# Synthesize of Ceo<sub>2</sub> Nanoparticles and Investigation of Antibacterial, Antifungal, Anticancer Activity

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# Abstract

The present research investigates the characterization and biological applications such as anti-bacterial, antifungal, the anticancer activity of synthesized  $CeO_2$  nanoparticles (NPs) by using the Microwave irradiation technique. The structural conformation and size distribution of  $CeO_2$  in the nanometer range has been studied by PXRD and TEM analysis. Further, the optical absorption (bandgap) and Functional group  $CeO_2$  NPs were determined by UV-Vis and FTIR respectively. Powder X-ray diffraction (PXRD) spectrum of synthesized  $CeO_2$  NPs was well- matched with JCPDS number 81-0792 which confirms the arrangement of atoms with a body- centered cubic structure along with cell parameters having average crystalline sizes on 49 nm. The  $CeO_2$  NPs sample containing the main elements such as Cerium metal and Oxygen, further astrong bond between cerium metal and oxygen has been confirmed by EDX and FTIR analysis respectively. Besides, the microbial studies, namely, anti-bacterial, anti-fungal and the response in human breast cancer cells exposed to  $CeO_2$  NPs were analyzed with MDA MB 231 by MTT assay.

Keywords: MDA-MB-231; MTT assay; Anti-bacterial; Antifungal

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# Introduction

# Experimental

#### Preparation of Cerium Oxide (CeO<sub>2</sub>) Nanoparticles

The term Nanotechnology is applied to the various concepts such as fabrication, characterization, explored comprehensively for multifunctional behavior and technological application of nanomaterial's sized and more overplay an important role in controlling the size shape of the properties of the material. Since fabricated nanomaterials have enormous uses in many fields [1-5] worldwide, due to their unique size and shape [6,7], this allows the large development in the Nanotechnological world.

Commercially a large volume of  $CeO_2$  NPs are fabricated because they act as a cracking catalyst in petroleum products, flash memory in electronic devices, gas leakage detecting sensors, polishing chemicals to polish mirrors in many electronic devices and wafers, andadditive fuel cells in diesel to increase combustion efficiency or mileage ((HEI, 2001; Park et al.,2008a). Since CeO<sub>2</sub> NP donates or stores oxygen which is based on the ecological system. If the environment is rich with oxygen, then CeO<sub>2</sub> itself acts as a catalyst but in the case of hydrocarbon combustions, it donates oxygen.

Among the many metal oxide NPs, the synthesized CeO, provides beneficial properties such as attractive morphology, highly reactive and strongly oxidizing agents. Since nanoparticlescan bind with the animal's cancer cell line, further these bindings prevent the growth of cancer cells and therefore nanoparticles have the major application of cancer prevention in the biological field [12-14]. The abnormal growth of cancer cells easily spreads to other parts of the body causing damage to organs for cancer patients. The available chemotherapeutic agents may provide the side effects and therefore it is necessary to prepare bio-compatible compounds in the place of drugs to cure cancer disease and further the cost of treatment should be less possible for cancer patients to cure cancer cells and to achieve these benefits, the nanoparticles have been introduced in this recent decade which provides the effective treatment for cancer patients. In this research, we show that the prepared CeO, NP is successfully active against the MDA-MB- 231 cell line[15-16]. Further, in this research, the important biological application such as the effective antimicrobial activity using the synthesized CeO<sub>2</sub> has been studied.

The Microwave irradiation technique was employed to synthesize  $CeO_2NPs$ . All reactants were purchased from Sigma Aldrich. The precursors used were 1g of Cerium (III) nitrate hexahydrate and 2 g Urea are added in 50 ml of distilled water along with AR grade  $CeO_2NPs$  and mixed well; these powders are converted to disk-shaped pellets with the help of a hydraulic handheld and then this pallet sample was kept at 800°C for 30 minutes in a microwave generating furnace. This heating process on the pellet sample which contains  $CeO_2NPs$  allows for the reduction of  $CeO_2NPs$ . After half an hour of the heating process, the pellet sample (containing  $CeO_2NPs$ ) has been burnout to finish the process. These samples were ground for 1hrin an agate mortar to make it into fine powder possessing uniform particle size. The obtained material was subjected to different characterization to confirm the appropriateness of device application.

#### **Characterizations techniques**

Characterizing the CeO<sub>2</sub> NPs is very essential which provides the structural information (shape, diameter, and average size CeO<sub>2</sub>) and many other various predictions such as absorption (or emission) spectrum, bandgap, and functional groups of synthesized CeO<sub>2</sub> NP sample. These various aspects of information can be obtained by using various instruments such as X-ray, UV- Vis spectrometer, FTIR, PL, EDX, and TEM. Among many instruments, TEM alone uses the beam of the electron, to reveal the surface morphological information of the CeO<sub>2</sub> NP sample. Except for TEM, other all instruments use a particular range of electromagnetic radiation to reveal the crystalline nature of the system, CeO<sub>2</sub> NP size, bandgap and functional groups, and thepresence of chemical elements in the CeO<sub>3</sub> NPs.

#### **Test Microorganisms**

Aspergillus Mucor fungi was used for carrying out the antimicrobial activity studies. These microorganisms were grown for 3 days at 37 °C in Actinomyces Isolation Media (AIM) broth (Himedia, Mumbai, India). The sensitivity of these microorganisms to the reference antibiotics was checked using mycostatin as a positive control.

#### Antifungal Activity of the Sample

The samples of iron oxide, erbium oxide and iron oxide/ erbium oxide nanoparticles were loaded on Potato Dextrose agar plates at three different volumes (10, 20, and 30  $\mu$ l) and swabbed with fungi such as Aspergillus and Mucor. Antifungal activities of the samples were determined by well diffusion method on Potato Dextrose Agar (PDA) medium [17]. The PDA medium was composed of (gl<sup>-1</sup>) potato infusion-200, dextrose -20, and agar-15. The PDA medium waspoured into the Petriplate; and after solidification, the inoculum was spread on the PDA plates with sterile swab moistened with the fungal suspension. All the plates were incubated at 37 °C for 3 days and finally the inhibition zone was analysed. These fungi were grown in Actinomyces Isolation Media (AIM) broth (HIMEDIA Mumbai). Mycostain was used as the positive controlto check the sensitivity against antibiotics by the well diffusion method on PDA medium.

#### Antibacterial Activity of the Sample

The antibacterial activity of iron oxide, erbium oxide and iron oxide/erbium oxide nanoparticles were tested against Escherichia coli, and Bacillus sp using disc diffusion method. The iron oxide, erbium oxide and iron oxide/erbium oxide nanoparticles were prepared in appropriate concentration of 1 mg/ml with dimethylsulfoxide solution for this process. Then, the dispersed nanoparticles were impregnated to each sterile disc by using micropipette. After thatthe discs were kept on culture swapped Mueller Hinton Agar medium using sterile force and allowed to incubate for 24 hrs. The average zone of inhibition diameter was measured inmillimeter (mm).

#### Cell Culture and Cell Line Maintenance

The human breast cancer cell lines MDA MB-231 were obtained. Then, these cell lines were grown as a monolayer in Dulbecco's modified Eagle's medium (DMEM: Hi Media Laboratories, Mumbai, India), which was supplemented with 10% fetal bovine serum, 100 U/mLpenicillin, and 100  $\mu$ g/mL streptomycin (Hi Media Laboratories Mumbai, India) cells grown at 37°C in incubator under 5% CO<sub>2</sub> with high humidity [18-19].

# MTT Assay Method for Evaluation of Cell Viability and Cytotoxicity

The anticancer activity of samples on human breast cancer cell lines MDA MB-231 was determined by the MTT (3-

(4, 5-dimethyl thiazol-2yl)-2, 5-diphenyl tetrazolium bromide) assay [20-21]. These cells (1 × 105/well) were plated in 0.2 ml of the cells with concentration of 1 × 105 cells/ml. The plates were incubated for 24 hrs in 5% CO<sub>2</sub> incubator for cytotoxicity. After incubation, normal breast (MDA MB-231) cells were cultured in 1:1 mixture of dimethyl sulfoxide (DMSO). Then, they were added to each well and mixed well by micropipette [22]. The percentage of viable cells was visualized by the development of purple color due to the formation of formazan crystals. The suspension was transferred to the cuvette of a spectrophotometer and observed significant variance/instability in the optical density (OD).Measurements were performed and the concentration required for a 50% inhibition of viability (IC<sub>50</sub>) was determined and used for the bioassays.

# **Results and Discussion**

#### PXRD of Cerium oxide (CeO<sub>2</sub>) metal oxide nanoparticle:

The PXRD pattern of  $\text{CeO}_2$  NPs, thus contains the particle size information and crystalline nature, which was synthesized by Microwave irradiation technique were present in Figure 1a. The peaks in fig.1a are assigned to the cubic structure of  $\text{CeO}_2$  with lattice pointsa=b=c=5.412 Å and matched with the JCPDS No. 81-7092. Using Debye - Scherer's formula, the average  $\text{CeO}_2$  crystallite size is found as 49 nm[23]. Several Bragg reflections with 2 $\theta$  values 28.5°, 33°, 47.5°, and 56.4° are observed corresponding to (111), (200), (220), (311), (222) and (422) (Figure 1)... Further, the crystallinity nature, particle size distribution, and phase formation have been confirmed by HRTEM.

#### TEM

The electrons passing in a TEM instrument through the potential barrier of the synthesized sample afford the topography [Lin P-C 2014, Hinterdorfer P 2011] information of the CeO<sub>2</sub> NPs. In fig.2 (TEM image) can be observed from CeO<sub>2</sub> NPs structured size between 45.112 to 48.556 nm which gives an average crystallite of 46.8 nm (Table 2), and further these crystalline nature [24-25] of the CeO<sub>2</sub> NPs has been confirmed by SsAED pattern (Figure 2d). The results from the TEM image are agreed with the result of XRD data having crystal planes (111) and (200). **EDX Analysis** 

EDX analysis of CeO<sub>2</sub> NPs was shown in Figure 3, which confirms that the major chemical element such as Cerium and oxygen were present with strong absorption range and homogeneous distributions, further this spectrum proves that the cerium atom uniformly combined with the element oxygen which is summarized in Table.3.



Figure 1: The PXRD pattern of CeO<sub>2</sub> Nanoparticles

**Table 1:** The deduced magnetic parameters for  $CoCe_x Fe_{2,x}O_4$  (0.0  $\leq x \leq 0.10$ ) samples obtained from the M-H loops at 100 K

Samples		Cellparameters (Å)	d space(Å)	Volume (Å)³	Crystalline size (nm)	Structure	
C1		5.4	1.759	158	49.1	FCC	
ICPDS		D-Spacing (Å)		Cell Parameters (Å)		Cell Volume (Å) <sup>3</sup>	
samples	CardNo.	Calculations	JCPDS	Calculations	JCPDS	Calculations	JCPDS
CeO <sub>2</sub>	81-0792	1.7596	1.7609	5.408	5.412	158.34	158.55

# FTIR

FTIR spectrum of  $\text{CeO}_2$  NP is shown in Figure 4. The band at 425 cm<sup>-1</sup> shows the presence of metal oxide, thus predicts the presence of a strong bond and stretching between metal cerium andoxygen (i.e. O-Ce-O) [26]. The major peak at the point of 706 cm-1 corresponds to the stretchingvibration

of doubly coordinated oxygen due to corner shared element oxygen to two cerium atoms. The peak at the point of 1466 cm-1 is assigned to the C-N stretching vibration of urea which was added in the synthesis process. The peaks at the point of 1467 and 3428 cm-1 correspond to stretching vibration of c=c and O-H respectively.



Figure 2: TEM images of Cerium Oxide (CeO<sub>2</sub>) Nanoparticles

Area (nm)	Mean (nm)	Min (nm)	Max (nm)
48.556	86.527	10.288	251
47.663	71.1	26.583	253.647
46.005	85.089	21.012	145
45.112	79.468	16.884	121
Average = 46	5.8 nm	I	

Table 2: The size distributions of CeO<sub>2</sub> NPs

# **UV-Vis spectroscopy**

UV-Vis affords the bandgap as well as the stability of the CeO<sub>2</sub> NPs. The UV-Vis of CeO<sub>2</sub> NPs is present in fig. 5a and 5b. The optical property of CeO<sub>2</sub> NP allows them to absorb with a particular range of wavelengths of UV-Vis radiations and maximum absorption and cut- off wavelengths were found at 270 nm and 430 nm respectively. In Figure 5c, it was observed that the bandgap energy of CeO<sub>2</sub> NPs be around 3eV. This shows that the synthesized CeO<sub>2</sub> NPs could be useful in few medical applications [27,28].

#### Photoluminescence (PL) spectroscop

PL spectrum of  $CeO_2$  is useful to detect the high-quality crystalline nature and the fine structure of the sample. PL at ambient room temperature was implemented to reveal the optical absorption of Cerium Oxide (CeO<sub>2</sub>) nanoparticles. In Figure 6, CeO<sub>2</sub> NP shows the UV emissionand green excitations at 284 nm and 550 nm respectively. These results were also reported in previous research [29]. In common, the presence of oxygen



Figure 3: The EDX pattern of CeO<sub>2</sub> Nanoparticles

Table 3: The percentage of chemical elements in the synthesized CeO<sub>2</sub>.NPs sample

Spectrum: Spectrum 525-Ce						
Element Series		Net	unn. C (% .Wt)	Norm. C (% .Wt)	Atom. C (% .at)	Error (% .Sigma) (Wt 3)
Cerium	L-series	94064	89.31	89.31	48.83	26.90
Oxygen	K-series	13405	10.69	10.69	51.17	1.08
Total:	100.00	100.00	100.00			

vacancies in the sample creates defects, thus are responsible for the creation of broad peaks at larger wavelengths of light.

#### **Anti-Fungal activity**

The synthesized metal CeO<sub>2</sub> NPs provides fewer hazards

to the outside environment, but its toxic activity against the various fungi is higher [30-32], to find out the antifungal activity of CeO2 NPs, the disc diffusion technique has been carried out and results are shown in fig.7 and further summarized in Table 5.

The cell viability effect (or zone inhibition) of  $\text{CeO}_2$  nanoparticles on the fungi is discussed by explaining the two



Figure 4: FTIR spectrum of Cerium Oxide (CeO<sub>2</sub>)

Observed Band Position (cm) <sup>-1</sup>	Assignments
1311	C-N Stretching vibration
1647	C=C Stretching vibration
3428	O-H Stretching vibration
425, 706	Cerium Oxide

Table 4: The peak positions and corresponding functional groups of synthesized CeO, NPs

different mechanisms. First, the creation of  $H_2O_2$  around the CeO<sub>2</sub> NP leads to the possible hydrogen bond development between (OH group) cellulose content of fungi and oxygen atom which causes higher zone inhibition. In second, the release of Ce<sup>2+</sup> ion causes damage to the cell membrane which leads to the growth of higher zone inhibition [34-45]. Results reveal that 300

mg of the CeO<sub>2</sub> NP sample provides a higher toxic activity than the toxic activity provided by the lower concentrations of CeO<sub>2</sub> NPs. The separate 300 mg of the CeO<sub>2</sub> NPs causes the higher zone inhibition as 20 mm and 15 mm respectively for *Mucor* and *Aspergillus*. Finally, the results from figure 7 show that the toxicity given by the CeO<sub>3</sub> is higher for *Mucor* than *Aspergillus*.



Figure 5a: UV-Vis absorbance spectra of CeO, NPs





9



Figure 5c: Photon energy vs.  $(\alpha h\nu)^2$  spectrum of CeO, NPs

Figure 7 the Anti-fungal activity of Crium oxide nanoparticles. 1) *Aspergillus* and 2) *Mucor* at 100 mg, 200 mg, and 300 mg, 3) concentration where the control.

#### Anti-cancer

The cytotoxicity of CeO, NPs with various concentrations (5  $\mu$ g, 10  $\mu$ g, 50  $\mu$ g, 75  $\mu$ g, and100  $\mu$ g /mL) against MDA-MB- 231 cell lines were investigated by the MTT method and its results were present in fig.9, further, the observed values are tabulated (Table 7). In the MTT assay, the cell morphology was captured after 48 hours to obtain the best results. In previous research, the significant results against MDA-MB-231 were observed by adding the 50 µg/mL or 75 µg/mL [52-55]. An advantage of CeO, NP is its available lesser size in the sample, so we believed that the synthesized lesser concentrations of CeO, NPs in this research, have the higher potential to destroy the MDA-MB-231 cell line. Therefore the chosen concentration of CeO<sub>2</sub> NPsin this research is lesser than the previously published results of the cytotoxicity study. The chosen lowest concentration  $(5 \,\mu g/ml)$  of CeO<sub>2</sub> NP provided the highest cell destruction with 86% cytotoxicity causing the lesser cell viability (14%.) of the MDA-MB-231 cell line.

# Conclusion

In this study, using Microwave radiation the CeO, NPs have been prepared and its effective anticancer and antimicrobial studies were successfully executed. The formed CeO<sub>2</sub> crystalline structure and its size range around 45 to 49 nm have been analyzed using XRD and TEM analysis. All the possible major elements and functional groups present in the sample which was synthesized by adding cerium nitrate, urea, and water molecules, have been successfully found using EDX and FTIR studies respectively. The bond between major element cerium and oxygen was identified by noting the absorbance peak at 706 cm-1 in the FTIR spectrum. Antifungal results show that the CeO, NP effectively kills higher the number of Mucor than Aspergillus, further the possible mechanism that causes damage to the fungi by using CeO<sub>2</sub> NP was discussed. The synthesized CeO, NPs were also found effective against BACILLUS SP and E. COLI bacteria. The results of the MTT assay using the synthesized sample against MDA-MB 231 cell lines indicated the higher percentage of toxicity of CeO<sub>2</sub> NPs. Therefore this study was successfully performed towards biological applications.

#### Funding



**Figure 6:** PL spectra of CeO<sub>2</sub> NPs

Nanoparticle	Aspergilus % of Zone inhibition					
	Concentration			Control		
	100 mg	200 mg	300 mg	17		
	12 mm	13 mm	15 mm	17 mm		
	Mucor % of Zone inhibition					
CeO, Nanoparticles	Concentration			Control		
-	100 mg	200 mg	300 mg	22 mm		
	14 mm 18 mm		20 mm	22 mm		

**Table 5:** Anti-Bactria of Cerium Oxide (CeO<sub>2</sub>)

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Figure 7: Anti-Fungal activity of Cerium Oxide (CeO)

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Nanoparticle	BACILLUS SP % of Zone inhibition				
	Concentration		Control		
	100 mg	200 mg	300 mg	10	
	14 mm	15 mm	17 mm	19 mm	
	E. COLI % of Zone inhibition				
CeO, Nanoparticles		Control			
-	100 mg	200 mg	300 mg	24	
	16 mm	20 mm	22 mm	24 MM	



Figure 8: Anti-Bactria of Cerium Oxide (CeO<sub>2</sub>)

Sample Particulars				
Description	Conc. (µg)	Cytotoxicity(%)	Cell viability(%)	Cytotoxic Re- activity
	5	86	14	Severe
C1	10	83	17	Severe
	50	76	24	Severe
	75	77	23	Severe
	100	80	20	Severe

Table 7: Anti-Bactria of Cerium Oxide (CeO,	,).
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# Cytotoxicity Direct Method Cell line: MDA MB-231Sample particulars: MC1



**Figure 9:** Anti-Bactria of Cerium Oxide (CeO<sub>2</sub>).







Figure 11: Anti-Bactria of Cerium Oxide (CeO<sub>2</sub>)

# **Conflict Of Interest**

The authors declare that they have no conflict of interest

# Availability of Data and Material

The authors confirm that the data supporting the findings of this research are available within thearticle and its supplementary materials.

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