

Synthesis, Characterization of $ZrO_2:Tb^{3+}$ (1-9 mol %) Nanophosphors for Blue Lighting Applications and Antibacterial Property

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Received: 2.09.2021; Revised: 20.10.2021; Accepted: 24.10.2021; Published: 24.11.2021

Abstract: This paper reports the structural, morphological, and antibacterial studies of $ZrO_2:Tb^{3+}$ nanophosphors (NPs). The $ZrO_2:Tb^{3+}$ NPs were synthesized by hydrothermal route using Amylamine as surfactant. $ZrO_2:Tb^{3+}$ nanophosphors was characterized by Powder X-ray Diffraction (PXRD), Scanning Electron Microscope (SEM), Diffuse reflectance spectroscopy (DRS), Photoluminescence (PL), Raman spectra, Fourier Transform Infrared radiation (FTIR) and Transmission Electron Microscope (TEM). PXRD analysis shows better crystallinity, cubic in-phase and good homogeneity of the synthesized phosphors were confirmed. When the Tb^{3+} concentration varies, we obtain blue emissions from $ZrO_2:Tb^{3+}$ NPs. $ZrO_2:Tb^{3+}$ NPs have a promising approach to blue light sources in the display application. SEM images show that $ZrO_2:Tb^{3+}$ nanophosphors have good morphology with a nonporous structure. TEM and SAED pattern confirms that $ZrO_2:Tb^{3+}$ nanophosphors are crystalline in nature. $ZrO_2:Tb^{3+}$ (9mol %) nanophosphors possessed a good antibacterial ability.

Keywords: nanophosphors; Raman spectra; hydrothermal technique; Rietveld refinement; surfactant.

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1. Introduction

The significance of nanophosphors in the field of research has been hugely developed because of their optical property [1]. They were broadly utilized in display applications [2], luminescent security ink [3], solar cell applications [4], bioimaging [5], photonic applications [6], etc. Zirconia assumes an essential job in the field of catalyst because it has a high surface-to-volume ratio [7]. Zirconium Oxide is used in the treatment of organic pollutants [8], biomedical applications [9], neuroscience [10], mammary cancer imaging in mice [11]. ZrO_2 is nanophosphors exists in three crystalline phases at different temperature, i.e., cubic phase (>2640K), tetragonal phase (1440-2640K) and monoclinic phase (<1440K) [12]. To date, different methodologies have been used to synthesize the ZrO_2 nanophosphors, i.e., coprecipitation method [13], Microwave method [14], Sol-gel method [15], spray pyrolysis deposition [16], and combustion method [17], etc.

In this work, we are using the hydrothermal method to synthesize $\text{ZrO}_2:\text{Tb}^{3+}$ nanophosphors because of high reaction rates and control over crystalline size and shape [10]. Amylamine as a surfactant is used to synthesize $\text{ZrO}_2:\text{Tb}^{3+}$ nanophosphors, and also it helps in stabilizing the nanoparticles [18]. The effect of Tb doping on the antibacterial properties of the Tb-doped ZrO_2 nanoparticles was investigated.

White light-emitting diodes are new-age lighting frameworks due to their novel properties like high productivity, low power utilization, and not containing harmful materials like mercury. Tb^{3+} doped materials show blue emission and green emission. As a result lot of attention must be put on Tb^{3+} doped hosts for their conceivable application in WLEDs. Moreover, there is a developing interest in growing profoundly effective rare earth (RE) doped ZrO_2 nanophosphors to deliver WLEDs.

Zirconia particles and Zirconia-based compounds are profoundly poisonous to microorganisms because of their mechanical properties and biocompatibility. Zirconia-based nonmaterial as antibacterial part have been utilized in dental earthenware and their covering of clinical gadgets [19]. The present investigation is committed to studying the phase, energy band gap, morphology, and antibacterial activity [20] of $\text{ZrO}_2:\text{Tb}^{3+}$ nanophosphors. The novelty of the present work is the ability to regulate the blue color by changing the concentration of Tb^{3+} .

2. Materials and Methods

2.1. Synthesis.

Hydrothermal method is used to synthesis $\text{ZrO}_2:\text{Tb}^{3+}$ nanophosphors. Zirconium (IV) propoxide and terbium (III) nitrate pentahydrate is a precursor required for synthesis. Zirconium (IV) propoxide was dissolved in methoxy ethanol and stirred continuously for 4 hrs. The nitrate salt of Terbium (III) Nitrate Pentahydrate is dissolved in double distilled water to get the aqueous solution of Terbium nitrate [21]. The reaction mixture is constantly stirred for 20min on a magnetic stirrer and mixed in zirconium propoxide solution and the reaction mixture's pH solution. Then, at that time, the pH of the solution was acclimatized to 10 by the addition of 2M of NaOH solution under constant stirring. The resulting reaction mixture was added to a Teflon-lined stainless steel autoclave of 300 ml, filled with 80% solvent constituted by 60 % double distilled water and 20% Amylamine, which acted as a surfactant [22]. Autoclave acts as temperature gradient was kept in a hot air oven at 150°C for 24 h reaction time. Then the reaction mixture was centrifuged and dried in a vacuum oven at 80°C for 6 hrs. Then, at this point, the autoclave was placed in a hot air furnace at 150°C for 24 hours (optimized time). After hydrothermal treatment, REMI R-24 Centrifuge was used to centrifuge the arrangement Machine at 5500 rpm for 10 min and washed with twofold refined water to remove salt, which is delivered during the reaction. The centrifugation process was rehashed multiple times. Precipitation was dried in a vacuum oven at 80°C for 6 hrs to remove water content. The final product was calcined at 600°C for 4h [23].

2.2. Antibacterial activity.

To study the antibacterial properties, the materials used are $\text{ZrO}_2:\text{Tb}^{3+}$ (9mol%), Mueller–Hinton media, Petri plates, filter paper disc. From MTCC (Microbial Type Culture Collection) Chandigarh, India, following materials such as Two Gram-positive bacteria *Bacillus subtilis*, *Staphylococcus aureus*, and Two Gram-Negative bacteria *Escherichia coli*,

Salmonella typhi subcultured on nutrient Mueller-Hinton and incubated at 37°C. The bacterial suspensions were prepared by selecting a single colony and cultured in different test tubes containing nutrient media for 24 h at 37°C.

To identify the potentiality of all nanoparticles in different concentrations range from 0.0025 µg/mL to 0.015 µg/mL as antibacterial agent, the bacterial strains were inoculated in sterile test tubes containing M-H media for 24 h at 37 °C. The bacterial growth was measured at a wavelength of 600 nm using a spectrophotometer; the ZrO₂:Tb³⁺ (9mol%) culture showed less growth compared to other different doped concentrations of nanoparticles. The synthesized ZrO₂:Tb³⁺ (9mol%) nanoparticle was used for further characterization.

Mueller–Hinton agar plates were prepared, and the entire surface of the plate inoculum was evenly spread. Before applying nanoparticles to the disc, let the plate dry. Different concentrations of nanoparticles were used for the antibacterial activity.

The agar plate was divided into six sections to understand the differential antibacterial activity by filter paper disc diffusion method using the suspension of bacteria spread on a Mueller–Hinton agar plate.

The different concentrations of synthesized ZrO₂:Tb³⁺ (9mol%) nanoparticles were dissolved in sterile water and briefly sonicated.

On the surface of the filter paper disc, each sample of the nanoparticles was loaded. The antimicrobial activity was surveyed by assessing the zone of inhibition against the test organisms with a caliper. The antimicrobial test for every bacterium made three steps [24].

3. Results and Discussion

3.1. Powder X-ray diffraction and Rietveld refinement.

The PXRD patterns of ZrO₂:Tb³⁺ (1–9mol %)NPs are shown in fig.1. The PXRD patterns show the cubic phase structure, and peaks are matched with standard JCPDS card No. 899069 for confirmation [25].

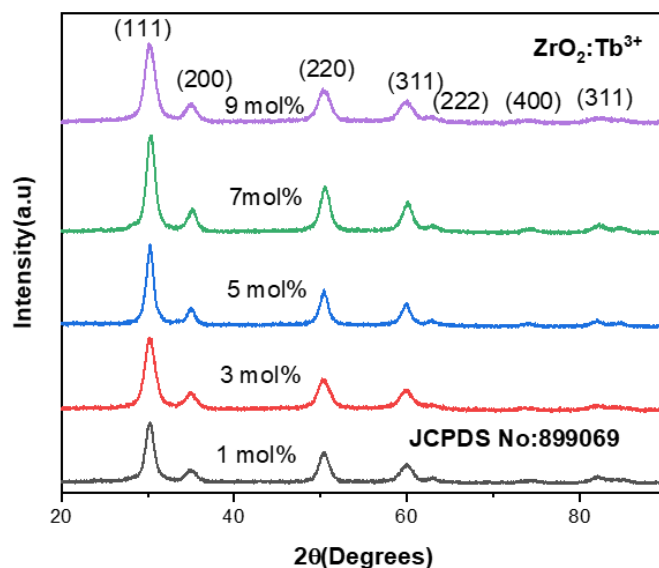


Figure 1. PXRD patterns of ZrO₂:Tb³⁺ (1–9mol %) nanophosphor.

Fig 2 shows Rietveld refinement of XRD patterns has been carried out using Fullprof suite software to provide proof for cubic phase with space group Fm-3m, which is in good agreement with JCPDS No. 899069. Cations have been set to (0, 0, 0) wyckoff from:4a, and O

atoms have been set to 1/4, 1/4, 1/4 wyckoff position:8c. In Rietveld refinement [26], technique intensity extraction occurred using the least square method and different experimental methods, and the sample-dependent peak broadening effect is studied. Here iterative refinement procedure is essential to obtain the potential fit between calculated and observed crystal structure. Parameters such as atomic positions, anisotropic displacement parameter, occupations, chemical binding, and geometry characteristics of the molecule in the structure model were modified using the least square methods [22].

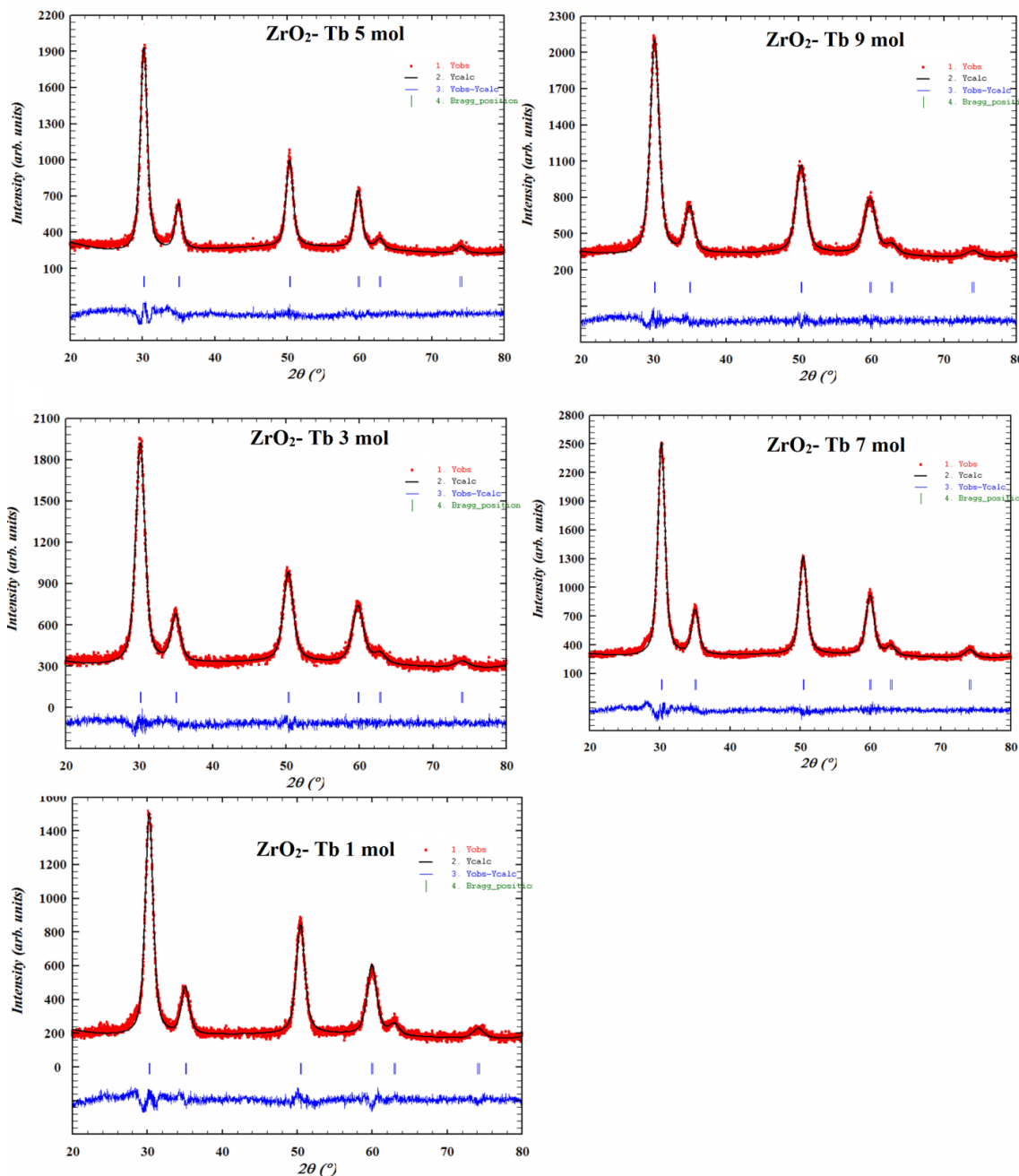


Figure 2. Rietveld refinement of ZrO₂:Tb³⁺ (1–9mol %) nanophosphors.

VESTA (Visualization for Electronic and Structural Analysis) software is used to study structural determination in crystallography, i.e., the volume of the unit cell, bond length, and angle [27]. Structural distortion in terms of bond length (R) and bond angle concerning dopant are tabulated in Table 1. There is no significant variation in the lattice parameters and volume

of the unit cell with increasing Tb^{3+} concentration [28]. Figure 3 shows the 3D structure of zirconia.

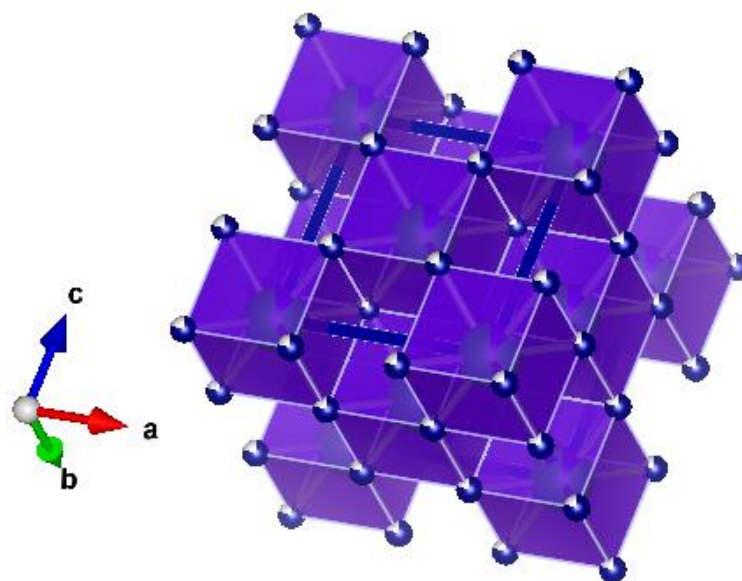


Figure 3. 3D structure of Zirconia.

Table 1. Rietveld Refinement Factors of $ZrO_2:Tb^{3+}$ (1–9mol %) nanophosphors.

Sr. No.	Sample	Goodness of fit (χ^2)	Lattice Parameter (Å)	Volume of Unit Cell (Å ³)	Zr-O(Å)	Zr-Zr(Å)	Electron Density around Zr	Oxygen Occupancy
1	$ZrO_2:Tb^{3+}$ (1 mol %)	1.49	5.1161	133.911	2.2153	3.6176	48.7022	1.8157
2	$ZrO_2:Tb^{3+}$ (3 mol %)	1.04	5.1279	134.839	2.2204	3.6260	48.2895	1.7776
3	$ZrO_2:Tb^{3+}$ (5 mol %)	1.86	5.1259	134.682	2.2196	3.6246	48.8062	1.6138
4	$ZrO_2:Tb^{3+}$ (7 mol %)	1.39	5.1172	133.997	2.2158	3.6184	48.5176	1.7346
5	$ZrO_2:Tb^{3+}$ (9 mol %)	1.06	5.1254	134.642	2.2194	3.6242	48.2535	1.6149

3.2. Scanning electron microscope.

Fig. 4(a-e) shows the SEM images of $ZrO_2:Tb^{3+}$ nanophosphors. It is observed that the morphology is uniform and similar to ovoid-like morphology. It has no voids and is porous.

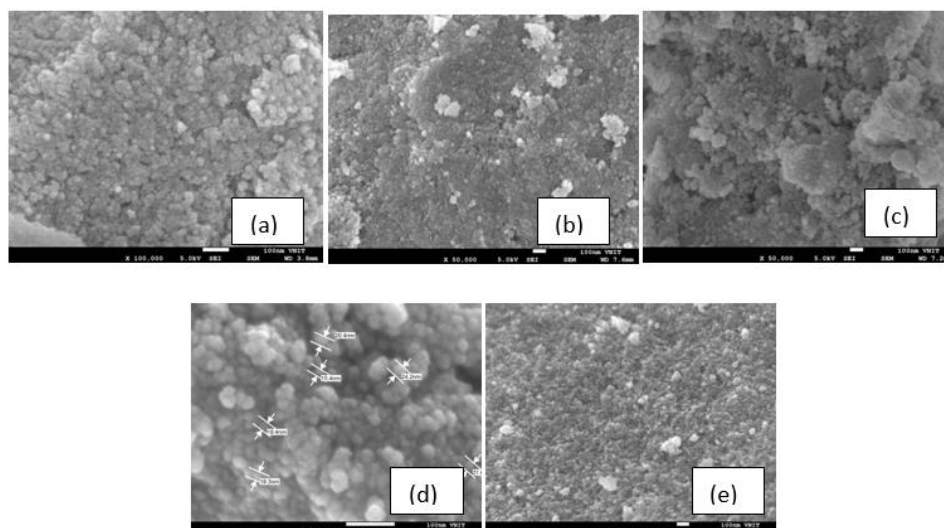


Figure 4. Scanning electron microscopy analysis of $ZrO_2:Tb^{3+}$ nanophosphors.

3.3. Diffuse reflectance spectroscopy.

Fig 5 shows the energy band gaps of $\text{ZrO}_2:\text{Tb}^{3+}$ (1-9mol%) nanophosphors. To calculate the bandgap of the NPs, Kubelka-Munk relation was used [29]. The bandgap of energy varies from 4.71–4.9 eV.

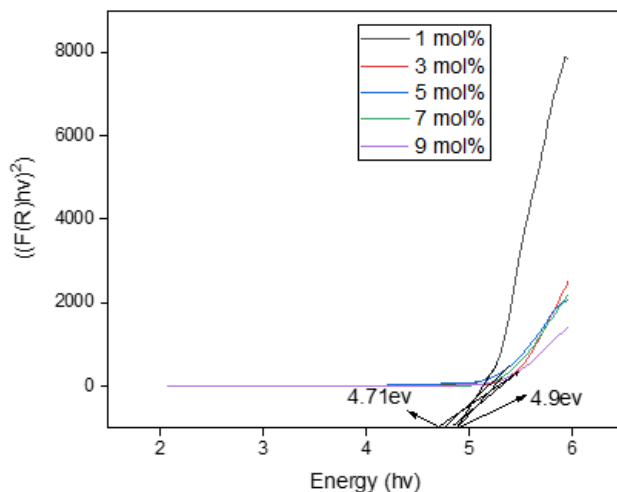


Figure 5. Energy band gaps of $\text{ZrO}_2:\text{Tb}^{3+}$ (1–9 mol %) nanophosphor.

3.4. Photoluminescence.

After excitation at 300 nm, emission spectrums were recorded for all nanophosphors, as shown in Fig. 6. The green emission above 480 nm was reported as a result of the advance of $^5\text{D}_4 \rightarrow ^7\text{F}_j$ transition, while the transition of $^5\text{D}_3 \rightarrow ^7\text{F}_j$ was responsible for the blue emission below 480 nm. In the present work, emission peaks at 466nm, 542nm because of $^5\text{D}_3 \rightarrow ^7\text{F}_5$, $^5\text{D}_4 \rightarrow ^7\text{F}_5$ transitions[1].

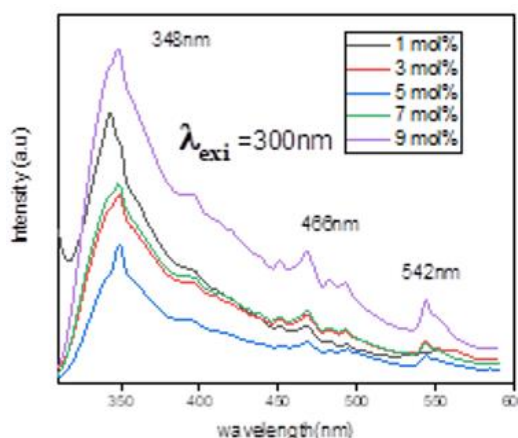


Figure 6. PL emission spectra of $\text{ZrO}_2:\text{Tb}^{3+}$ (1–9 mol%) nanophosphor excited at 300 nm.

The luminescent emission color of $\text{ZrO}_2:\text{Tb}^{3+}$ (1-9 mol%) NPs indicated by the 1931 chromaticity coordinates of the International Commission on Illumination (CIE). These coordinates represent the distinctive visible colors produced by the phosphors. The CIE chromaticity coordinates for the optimized sample were calculated from the emission spectra measured under 300 nm excitation wavelength, and the results are shown in Fig.7[30]. When the Tb^{3+} concentration varied, the color coordinates in the chromaticity diagram ranged from deep blue to light blue[31]. Table 2 shows the variation of Commission International de

L'Eclairage (CIE) chromaticity coordinates of the $\text{ZrO}_2:\text{Tb}^{3+}$ (1–9 mol%) phosphors under excitation wavelength at 300 nm.

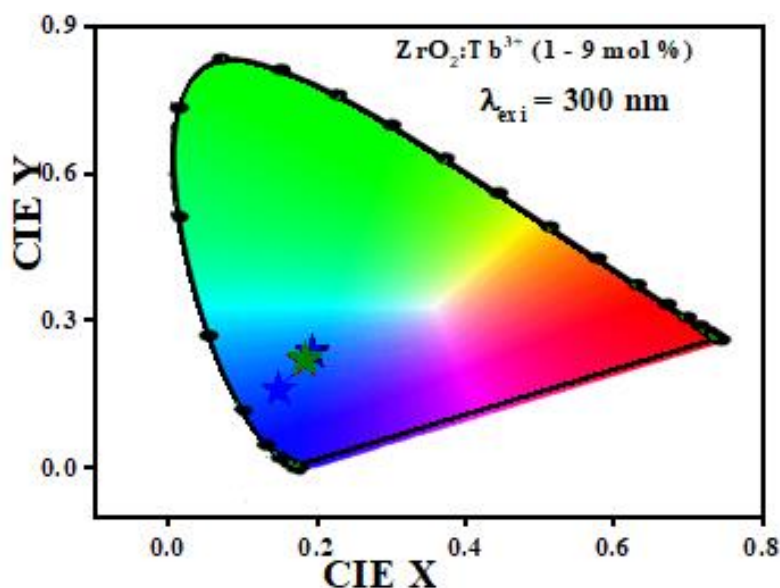


Figure 7. CIE chromaticity of $\text{ZrO}_2:\text{Tb}^{3+}$ (1–9 mol%) nanophosphors.

Table 2. CIE chromaticity coordinates of the $\text{ZrO}_2:\text{Tb}^{3+}$ (1–9 mol%) nanophosphors under excitation wavelength at 300 nm.

mol%	x	y
1	0.148172	0.159479
3	0.19355	0.235565
5	0.190875	0.239043
7	0.183765	0.218761
9	0.184044	0.220916

3.5. Raman spectra.

The Raman spectra of $\text{ZrO}_2:\text{Tb}^{3+}$ (1–9 mol %) NPs recorded in the range of 530–1000 cm^{-1} are shown in Figure 8. From Figure 8, the peaks were observed at 548 cm^{-1} (A_{1g}), 563 cm^{-1} (A_{1g}) [25], and 651 cm^{-1} (T_{2g}) [32]. In Figure 8, the peaks correspond to the cubic $\text{ZrO}_2:\text{Tb}^{3+}$ (1–9 mol%) NPs.

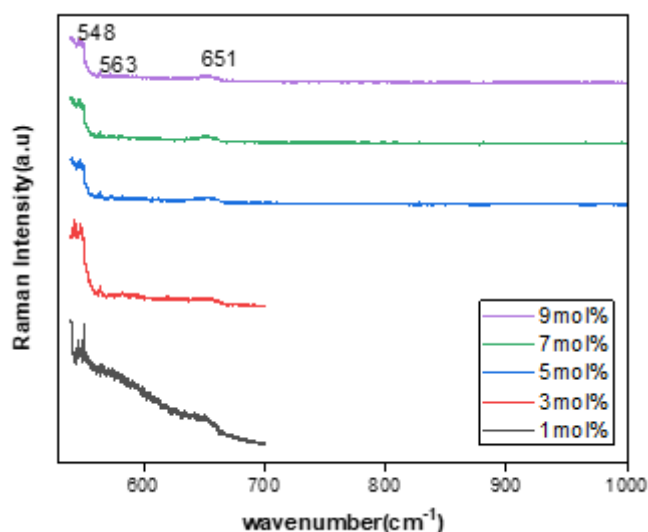


Figure 8. Raman spectra of the $\text{ZrO}_2:\text{Tb}^{3+}$ (1–9 mol %) NPs.

3.6. Fourier Transform Infrared radiation.

Figure 9 shows the FTIR spectra of the $\text{ZrO}_2:\text{Tb}^{3+}$ (1-9 mol %) Nanophosphors recorded in the range from 450 to 4500cm^{-1} . It gives information about the absolute OH content of the phosphors. The stretching frequencies of Zr – O bond is due to the band at 1133 cm^{-1} . The bending vibration of the OH group is due to 1424 cm^{-1} . The bonds at 3664 cm^{-1} due to stretching vibrations of absorbed water molecules in the form of OH group vibrations also show low absorbance [33]. The peak at 1715cm^{-1} is due to the characteristic of the vibration of OH groups. The characteristic of the vibration of OH groups is due to 1715cm^{-1} [34].

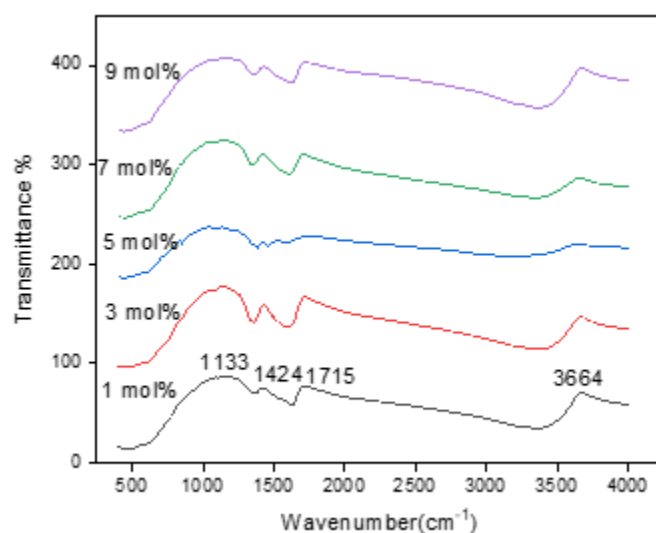


Figure 9. FTIR spectra of the $\text{ZrO}_2:\text{Tb}^{3+}$ (1-9 mol %) NPs.

3.6. Transmission electron microscope.

Fig 10 shows the TEM and SAED patterns of $\text{ZrO}_2:\text{Tb}^{3+}$ (9mol %) nanophosphors. Tem images show particles are tightly packed. SAED pattern confirms crystalline structure.

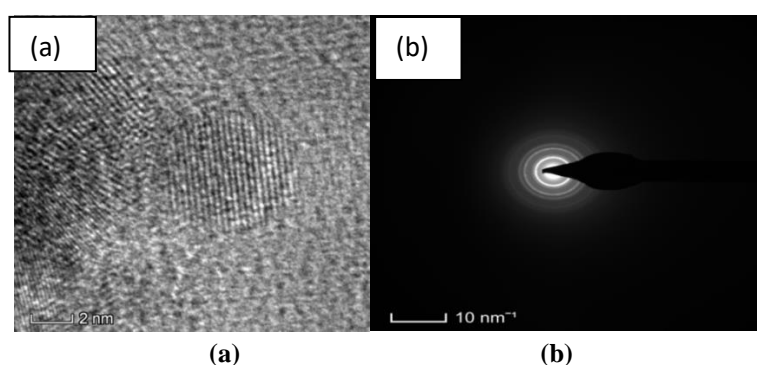


Figure 10. (a) TEM and (b) SAED pattern of $\text{ZrO}_2:\text{Tb}^{3+}$ (9 mol%) nanophosphors.

3.7. Antibacterial activity.

The filter paper disk diffusion assay was performed to test the inhibition zone of $\text{ZrO}_2:\text{Tb}^{3+}$ (9mol%) NPs against human pathogenic Gram-positive *B. subtilis*, *S. aureus*, and Gram-negative *E coli*, *S typhi* bacteria. Here, the $\text{ZrO}_2:\text{Tb}^{3+}$ (9mol%)NPs showed activity against all the tested bacteria at different concentrations. The data show that the inhibition zone diameter for synthesized $\text{ZrO}_2:\text{Tb}^{3+}$ (9mol%)NPs gradually increased with increasing

concentration. The antibacterial activities were tested against four bacteria species which include two Gram-positive bacteria: *B. subtilis*, *S. aureus* inhibition showed higher than the other two Gram-negative bacteria: *E. coli*, *S typhi*.

Table 3 shows the antibacterial activity studies. Increased inhibition of the zone leads to the destruction of bacterial cells due to the increased formation of reactive oxygen species (ROS) [35]. These elevated ROS disrupts the bacterial cell membrane by lipid peroxidation and lead to a high level of membrane leakage results in bacterial death with increasing concentration of nanoparticles [36]. In the present study, the binding of $ZrO_2:Tb^{3+}$ (9mol%) NPs to bacteria depends on the surface area available for interaction with the cell surface, which penetrates and changes take place in its cell membrane morphology and physiology. The ROS modulates cell function, which leads to DNA damage, protein denaturation, turnover of lipids induces oxidative stress. This will enable to generate ROS through different biochemical pathways. The DNA damage is due to induced ROS [37].

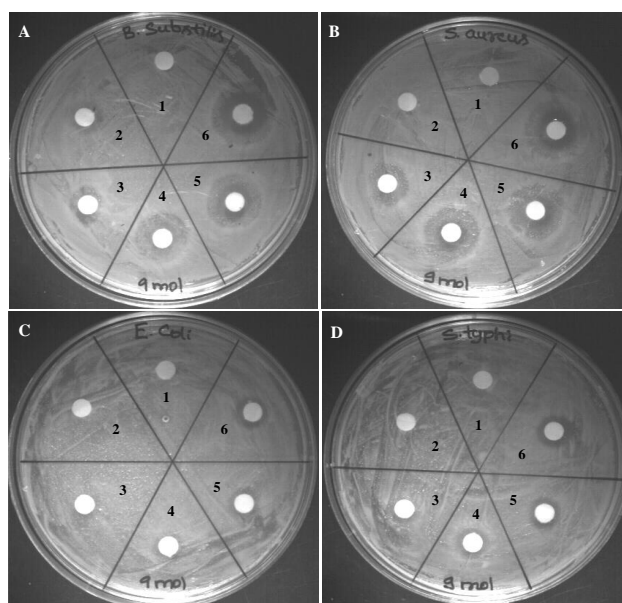


Figure 11. Antibacterial effects of $ZrO_2:Tb^{3+}$ (9mol%). 1. Control, 2. 0.0025 $\mu\text{g/mL}$, 3. 0.005 $\mu\text{g/mL}$, 4. 0.0075 $\mu\text{g/mL}$, 5. 0. 01 $\mu\text{g/mL}$ and 6. 0.015 $\mu\text{g/mL}$.

Table 3. Antibacterial activity of $ZrO_2:Tb^{3+}$ (9mol%) nanophosphors on Gram-positive; *Bacillus subtilis*, *Staphylococcus aureus*, and Gram-negative; *Escherichia coli*, *Salmonella typhi*.

$ZrO_2:Tb^{3+}$ (9mol%)(μg)	Gram-positive bacteria		Gram-negative bacteria	
	Zone of inhibition in mm			
	<i>B. subtilis</i>	<i>S. aureus</i>	<i>E. coli</i>	<i>S. typhi</i>
Control	0.00	0.0	0.00	0.00
25	0.13	0.14	0.00	0.00
50	0.24	0.27	0.00	0.11
75	0.35	0.36	0.21	0.23
100	0.43	0.45	0.33	0.32
150	0.52	0.58	0.42	0.41

4. Conclusions

Tb doped zirconium oxide NPs were synthesized by Hydrothermal synthesis route with Amylamineas surfactant. PXRD image reveals it is cubic in shape. Retevield refinement confirms cubic in phase; Raman patterns support PXRD patterns. DRS pattern shows bandgap varies from 4.17-4.9ev. SEM and TEM provide morphology of $ZrO_2:Tb^{3+}$ NPs. SAED pattern confirms crystalline in phase. The $ZrO_2:Tb^{3+}$ NPs belong to one kind of novel optical material

as a promising future candidate for strong blue emission and have drawn an increasing amount of attention. In the antibacterial activity of ZrO₂:Tb³⁺ (9mol%), NPs were more effective against selected Gram-positive bacteria than on selected Gram-negative bacteria.

Funding

This research received no external funding.

Acknowledgments

The authors are grateful to Prof. Smita A Acharya, Rashtrasant Tukadoji Maharaj Nagpur University, IITM SAIF, INUP-IISc, and CeNSE, Bangalore, for the characterization of the compound.

Conflicts of Interest

The authors declare no conflict of interest.

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