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Keywords: CeO₂, thin film, radiation shielding, Sol-Gel

Abstract

In this study, CeO₂ thin films were produced using the spin coating method, which is one of the sol-gel methods, in six different molarities. X-ray diffraction (XRD) patterns revealed the characteristic peaks of the films, while Field Emission Scanning Electron Microscopy (FESEM) confirmed their homogeneous structure. Then, radiation shielding parameters like linear absorption coefficient (LAC), mass absorption coefficient (MAC), tenth value layer (TVL), mean free path (MFP), and half value layer (HVL) were thoroughly examined. The results showed that increasing molarity had a significant effect on the thickness values of thin films and the absorption parameters were found to improve with increasing molarity. Both LAC and MAC values decrease as the energy level increases, but the increase in CeO₂ molarity leads to a strong increase on these coefficients. The HVL value was also found to be 0.42 cm at the lowest energy of 14.957 keV and to be around 10 cm at the greatest energy of 59.543 keV (0.05 M). When the radiation energy applied to the material was raised from 14.957 keV to 59.543 keV, it was found that the MFP values of 0.05 M CeO₂ thin films grew gradually from 0.61 cm to 14.51 cm. High energy radiation of 59.543 keV and a low density (0.05 M) medium resulted in peak TVL values of 33.423 cm, allowing the radiation to pass through the material with minimal interaction.

1. Introduction

As a result of the advancement of technology and its widespread use in industry, people are exposed to more ionizing radiation [1]. Scientists have focused on various studies to protect people from the harmful effects of radiation. One of these studies is on the use of thin film technology as a radiation shielding. The use of thin films as radiation shields is not only important for human health but also for technological devices [2–4]. Dense materials such as lead and concrete are generally used as radiation shielding [5]. However, the low optical transmittance of lead poses a major obstacle to its use in some applications of technology. For example, since lead is used in the production of glasses used for radiation protection, the optical transmittance of these glasses is quite low. This reduces the quality of vision when using these glasses. In the literature research, it has been observed that optical transmittance decreases as the lead doping ratio increases in lead doped thin films [6, 7]. Fabrication of thin films with high optical transmittance and radiation absorption capacity is important for human health and some technological applications. In this study, the radiation absorption effects of CeO₂ thin films produced in six different thicknesses were examined in detail depending on five different energies. CeO₂

thin films stand out as a fascinating material with numerous applications. CeO_2 possesses exceptional properties, including high atomic number, a wide band gap, high dielectric constant, high chemical stability, remarkable mechanical strength, excellent lattice compatibility, a good refractive index for optical applications and transparency in the visible and near-infrared spectral regions [8–10]. The high atomic number and density of these elements contribute significantly to their ability to attenuate and absorb ionizing radiation [11]. Considering these properties, it shows that CeO_2 thin films can be preferred as radiation shields. When literature studies are examined, it is seen that although studies investigating CeO_2 thin films as radiation shields are encountered, the studies conducted are limited. In addition, this study differs from other studies in the literature in terms of preparing thin films produced at different molarities in various thicknesses and examining the absorption properties depending on thickness and energy by exposing these films to radiation at six different energies. In recent years, CeO_2 thin films have been successfully utilized in various technological applications, including catalytic [12], electrochemical [13], optical [14], electrical [15], gas sensor [16], corrosion-resistant applications [17, 18] and UV-blocking filters [19]. In this study, homogeneous and economical films were obtained by using the spin coating method for the fabrication of CeO_2 thin films. One of the reasons for using the spin coating technique is that it is very easy to apply and can be obtained at a relatively low cost. The physical properties of the films obtained depend on thermal annealing and thermal annealing time [20–22]. However, various coating parameters such as coating speed, coating time and molarity of the precursor solution also play a role in the physical properties of thin films during coating [23, 24]. In this study, the radiation absorption parameters of CeO_2 thin films with six different thicknesses and five different energies were investigated in detail. Considering the large number of studies conducted in this field [25–27], it is thought that this study will make important contributions to the literature.

2. Experimental section

2.1. Methodology

The system is called a narrow geometry Am-241 radioisotope source attenuation geometry. The Americium-241 radioactive annular source was selected because to its compatibility with secondary targets, ability to facilitate non-destructive excitation, independence from a power source, and capacity to generate an adequate quantity of gamma rays. Photons from the Am-241 source, which produces photons with an energy of 59.5 keV, are sent to the detector without a sample. The number of photons is recorded as a blank count. Then, the sample is added, and the same process is performed. This time, the photons passing through the sample are recorded. According to the Lambert-Beer law, calculations are made with these photon numbers, and the mass absorption coefficient is calculated. Taking the sample density into account, the linear absorption coefficient and the mean free path, half-value thickness and tenth value thickness parameters derived from here are calculated.

2.2. Fabrication and characterization of CeO_2 thin films

CeO_2 thin films were coated on the c-Si surface by spin coating method. Cerium (III) chloride heptahydrate ($\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, Sigma-Aldrich) with a purity of 99% was used as the source of cerium, while ethyl alcohol was preferred as the solvent. 0.2 M citric acid monohydrate (Merck, 99.7%) was added to the solution, stirred in a magnetic stirrer for 30 min and aged for 24 h. To increase the adhesion of thin films, 75 μl of triethanolamine (97%) was added to 5 ml of solution and mixed at 50 °C for half an hour. The coating was carried out on CZ-Si p-type wafers with a thickness of 725 μm and a size of 1 cm \times 1 cm. The wafers were cleaned for 5 min in an ultrasonic bath using ethyl alcohol and deionized water. Then the natural SiO_2 layer was removed with HF (1:10) solution. The coating was carried out at a speed of 6000 rpm for 10 seconds so that the solution spreads over the surface and dried at 200 °C for 5 min. XRD patterns were obtained using a Rigaku Smartlab diffractometer with $\text{CuK}\alpha$ radiation ($\lambda = 1.5408 \text{ \AA}$) to examine the crystal structures of the thin films. FESEM images were performed with the Zeiss Sigma 300 model to examine the surface morphology of thin films.

2.3. Gamma ray attenuation measurements

The experimental configuration is illustrated in figure 1. It encompasses the secondary source, the primary source (Am-241), the absorber, and the detecting apparatus. The secondary source beam strikes the Am target, while the absorber collects the emitted beam from the Am target. A collimated Ultra-LEGe (Ortec) detector was used to measure the transmitted gamma rays. The detector exhibits a relative yield of 25% at 1.33 MeV and a resolution of 1.85 keV at the same energy level. The output preamplifier, capable of rejecting pulse pile-ups, transmitted signals to a multichannel analyser linked to a personal computer equipped with appropriate software for data collection and peak analysis. A peak fitting program conducted the fitting and deconvolution procedures. Radiation absorption measurements at different energies were performed to determine the shielding parameters of thin films.

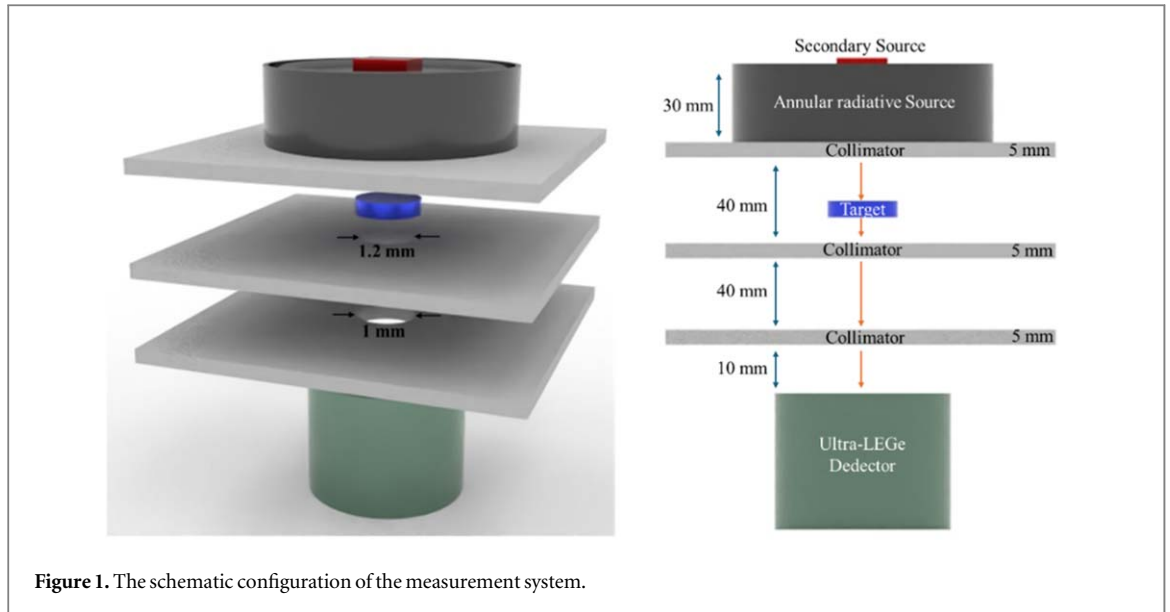


Figure 1. The schematic configuration of the measurement system.

2.4. Calculations

The attenuation coefficients were determined using the Lambert–Beer equation for Cerium dioxide thin films at various molarity levels, employing prior analytical results [28];

$$I = I_0 e^{-\mu x} = I_0 e^{-\left(\frac{\mu}{\rho}\right) d} \quad (1)$$

I_0 represents the incident beam directed towards the detector from the x-ray source in the absence of any sample. The beam is conveyed from the sample to the detector. d represents the mass per unit area (g/cm^2), whereas μ/ρ denotes the mass attenuation coefficient (cm^2/g) [29];

$$\left(\frac{\mu}{\rho}\right)_c = \sum_i W_i (\mu/\rho)_i \quad (2)$$

where W_i represents the weight fraction and the mass attenuation coefficient of the i th constituent element, respectively. The weight fraction of a chemical substance is expressed by equation (3) [30];

$$W_i = \frac{n_i A_i}{\sum_i n_i A_i} \quad (3)$$

The atomic weight of the i th element is denoted as A_i , while the quantity of formula units is denoted as n_i .

Theoretical values of the MAC (cm) were acquired from the XCOM database [31].

The LAC ($\mu \text{ cm}^{-1}$) quantifies the rate of photon absorption per unit thickness. This is determined by multiplying the mass attenuation coefficient by the specific gravity of the substance and is derived by multiplying the MAC by the material's specific gravity, as outlined below [32];

$$\mu = \mu_m \rho \quad (4)$$

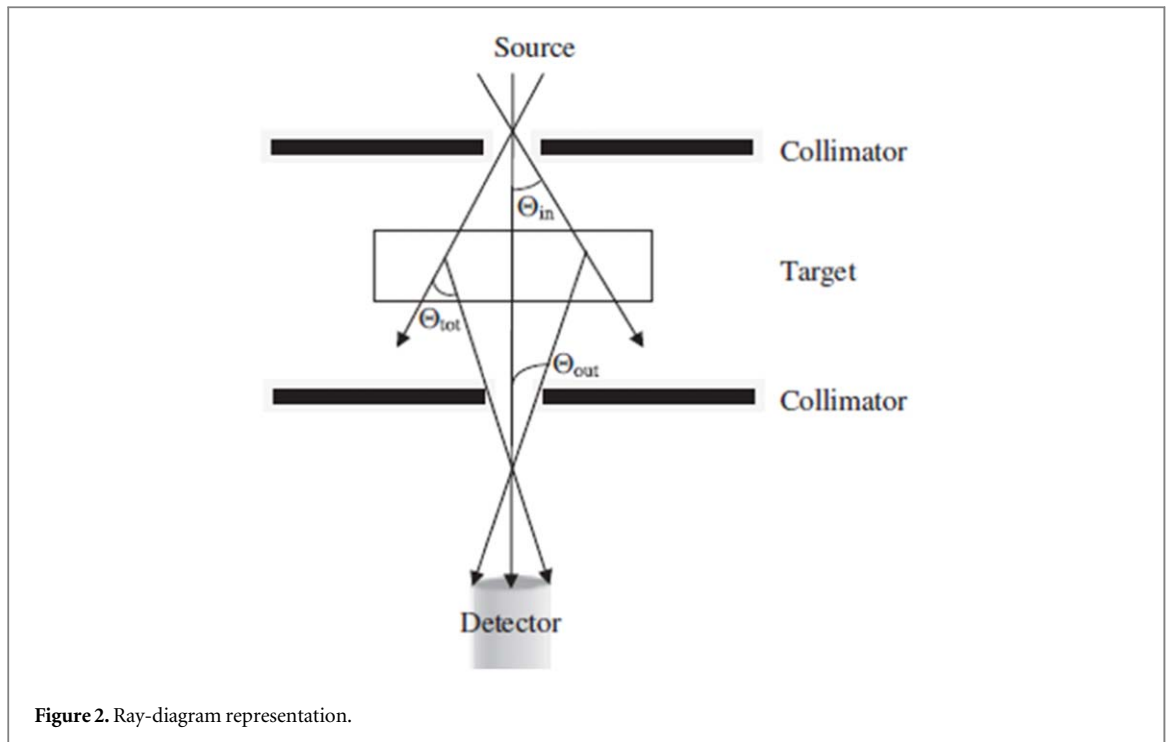
where μ_m represents the mass attenuation coefficient of the material and ρ denotes the density of the substance. The MFP (cm) is a crucial metric that indicates radiation attenuation. The MFP in physics refers to the average distance a moving particle (such as an atom, molecule, or photon) travels before substantially changing its direction, energy, or other attributes in a certain setting. This typically results from one or more subsequent collisions with other particles. The mean free path values are determined using the subsequent equation: [33];

$$MFP = (1/\mu) \quad (5)$$

The substance retains residual energy following the establishment of atomic binding energy or secondary photons. The thickness of a material, referred to as the HVL (cm), functions to diminish photon intensity and assess its capacity to attenuate gamma rays. The half-value layer (HVL), or half-value thickness, is the thickness at which the intensity of radiation passing through a material is reduced by fifty percent. The subsequent calculation is employed to ascertain the HVL [33];

$$HVL = \ln 2 / \mu \quad (6)$$

TVL, or the tenth-value layer, is a comparable concept. The entire amount of material necessary to attenuate 90% of the radiation, so reducing it to one-tenth of its original intensity, is referred to as the TVL, as indicated in equation (5) [33];



$$TVL = \ln 10 / \mu \quad (7)$$

where μ is the total LAC.

This technique facilitates the calculation of gamma-ray absorption parameters of CeO_2 thin films of six different molarities by using narrow-beam x-ray transmission geometry to measure different gamma energies. Figure 1 illustrates the experimental configuration, comprising the secondary source, primary source americium-241 (Am-241), absorber, and detection system. The Am-241 radioactive annular source was selected because to its compatibility with secondary targets, ability to facilitate non-destructive excitation, independence from a power source, and capacity to generate an adequate quantity of gamma rays. The secondary source beam impacts the Am target, while the absorber captures the emitted beam from the Am-241 target.

Lead collimators around the detector to obstruct radiation from adjacent objects from accessing it. Three collimators with apertures of 1.2, 1.2, and 1 mm were placed at distances of 30, 75, and 120 mm from the secondary source, respectively. The width of each collimator measured 5 mm. The samples were placed on a specimen holder, 70 mm from the secondary source. The separation between the secondary source and the detector is 135 mm [34]. Figure 2 illustrates the total scatter acceptance angle (θ_{tot}), defined as the sum of the angle encompassed by the exit collimation (θ_{out}) and the angle represented by the incident beam divergence (θ_{in}).

3. Result and discussion

XRD measurements were carried out in the range of 25° to 50° to perform the structure analysis of thin films produced at different molarities. As seen in figure 3(a), three peaks belonging to CeO_2 thin films were observed and the main peak was determined to be in the (111) plane at 28.33° . It was observed that the peaks of CeO_2 thin films were consistent with the literature and the peak intensity increased as the thin film thickness increased [35, 36]. Considering that D is the crystal size, λ is the x-ray wavelength, θ is the Bragg angle, and β is the half-peak width, the average crystal size of thin films can be easily calculated by the Debye–Scherrer equation (equation (8)) based on the x-ray diffraction patterns [37, 38].

$$D = 0.9\lambda / \beta \cos \theta \quad (8)$$

The crystal size results calculated by taking into account the (111) plane, which is the main peak of CeO_2 thin films, are given in figure 3(b). According to the results obtained, the average crystal size increases with increasing molarity [39].

In figure 3(b), the average crystal size values of thin films prepared at five different molarities are presented. Since no CeO_2 peak was observed in the 0.05 M solution, the crystal size could not be calculated. However, the crystal size of the thin film obtained from the 0.1 M solution was calculated as 5.80 nm. For the 0.2 M, 0.3 M, 0.4 M, and 0.5 M solutions, the crystal sizes were determined to be 7.40 nm, 9.10 nm, 10.40 nm, and 10.10 nm,

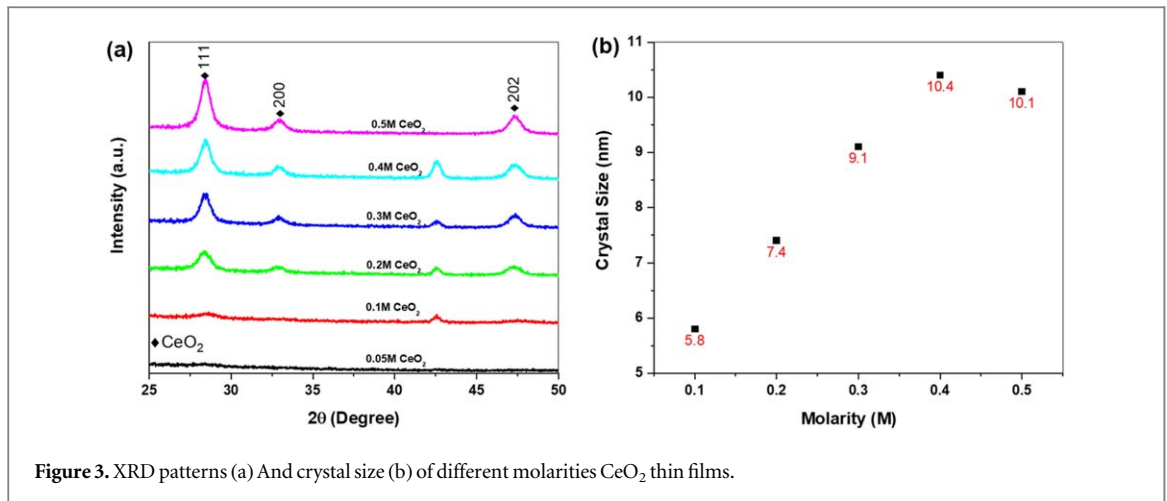


Figure 3. XRD patterns (a) And crystal size (b) of different molarities CeO_2 thin films.

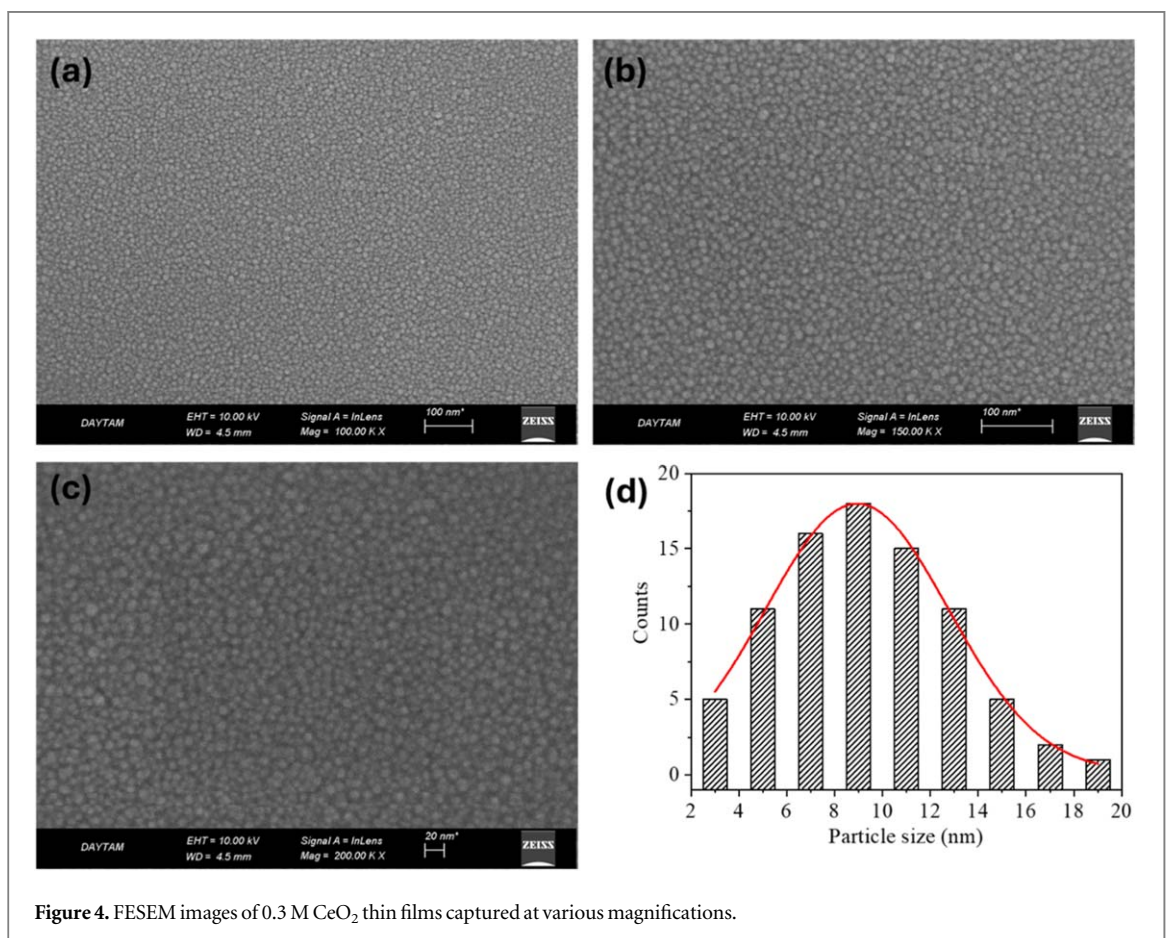


Figure 4. FESEM images of 0.3 M CeO_2 thin films captured at various magnifications.

respectively. The average crystal size for the (111) plane of the thin films prepared at these five molarities was calculated as 8.56 nm, which is consistent with the literature [40]. Figure 4 shows the results of FESEM investigations at various magnifications to investigate the surface morphology of 0.3M CeO_2 thin films. The obtained results show that the CeO_2 thin film has a granular structure and is uniformly dispersed over the surface of the crystalline silicon. FESEM pictures, in addition to XRD investigations, verify the homogenous film production and the structure of CeO_2 thin films. The particle size was determined from image analysis of FESEM observations in conjunction with ImageJ software. Histogram plot of the samples prepared at 0.3 M CeO_2 thin films is shown figure 4(d). The particle sizes of CeO_2 thin films were calculated as 8.93 nm, which is seen to be in good agreement with the value obtained from XRD measurements.

Also, The thickness values corresponding to each molarity for thin films prepared at different molarities are presented in figure 5. The thickness values were calculated theoretically by fitting the reflectance curves in previous study [41] and using the Fresnel equations in equation (9).

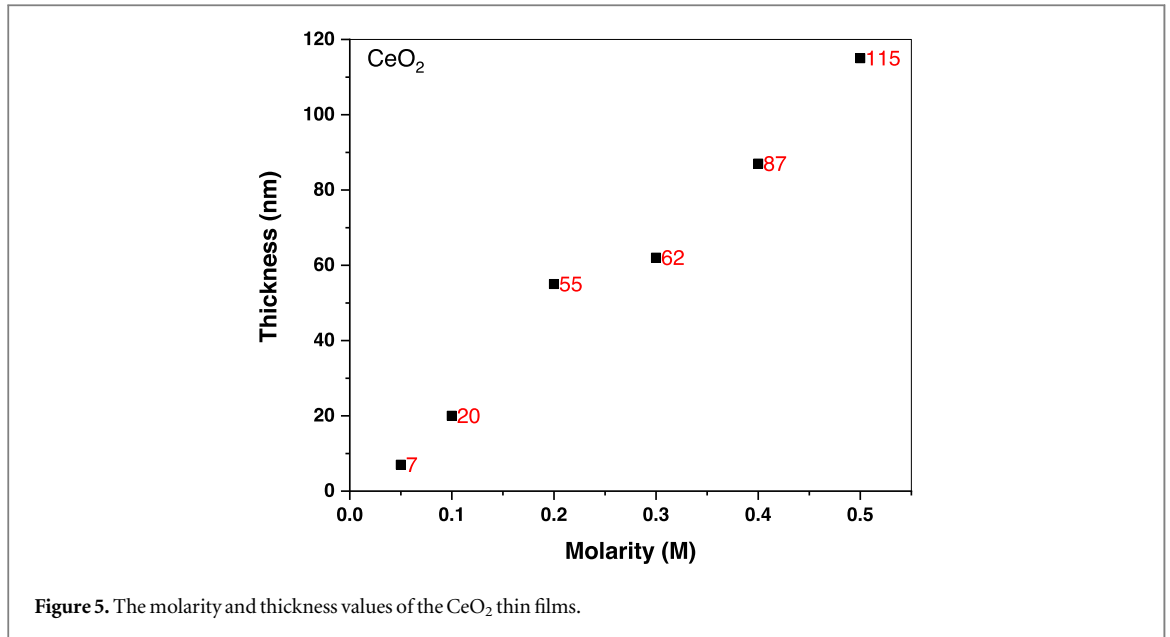


Figure 5. The molarity and thickness values of the CeO₂ thin films.

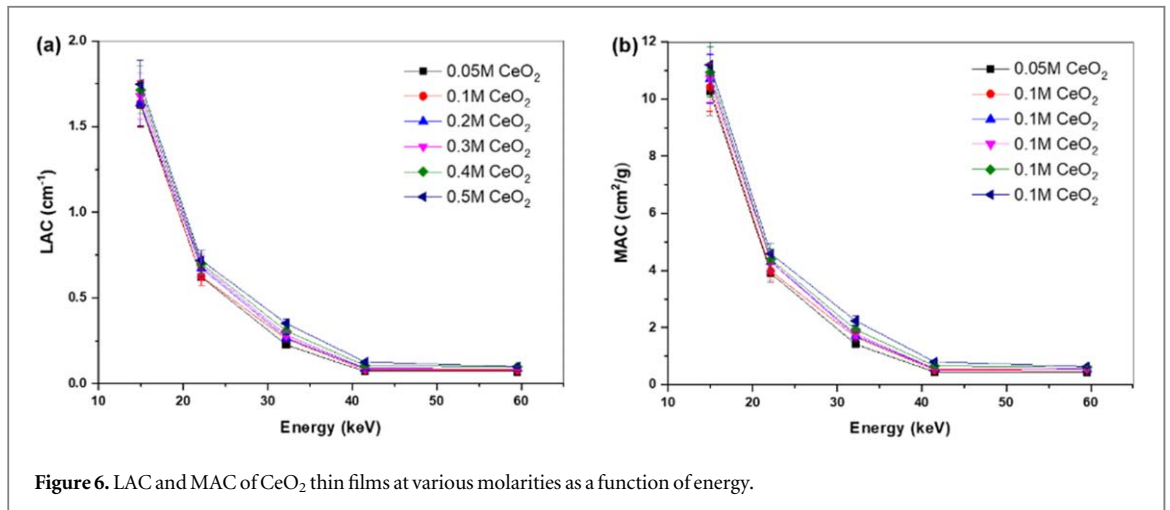


Figure 6. LAC and MAC of CeO₂ thin films at various molarities as a function of energy.

$$R = |r^2| = \frac{r_1^2 + r_2^2 + 2r_1r_2 \cos 2\theta}{1 + r_1^2r_2^2 + 2r_1r_2 \cos 2\theta} \quad (9)$$

where, $r_1 = \frac{n_0 - n_1}{n_0 + n_1}$, $r_2 = \frac{n_1 - n_2}{n_1 + n_2}$, $\theta = \frac{2\pi n_1 d_1}{\lambda}$, d_1 is thin film thickness, n_1 is thin film refractive index, n_0 is the air refractive index, n_2 is substrate refractive index, and θ is light incidence angle [42, 43].

As can be seen from the figure 5, increasing molarity caused an increase in thin film thickness. For example, while the 0.05 M CeO₂ thin films with the lowest molarity had a thickness of 7 nm, a thickness value of around 115 nm was obtained for 0.5M CeO₂ thin films. In the following stages of the study, the thickness values corresponding to each molarity were used to calculate the density of the thin films and experimental calculations were carried out accordingly.

Determination of shielding parameters of thin films is very important in terms of determining the usage areas of the material. In this study, LAC, MAC, TLV, MFP and HVL parameters were determined experimentally as shielding parameters, respectively. Figure 6 and table 1 show the LAC (μ) and MAC (μ/ρ) values of samples with different CeO₂ concentrations at specific energy levels such as 14.957 keV, 22.162 keV, 32.191 keV, 41.529 keV and 59.543 keV. The results in the table reveal that both LAC and MAC values decrease as the energy level increases, but the increase in CeO₂ molarity leads to a strong increase on these coefficients. For example, at 14.957 keV energy, LAC is 2.685 for 0.05 M CeO₂, while this value increases to 7.239 at 0.5 M CeO₂. Similarly, at 0.05 M CeO₂, MAC is 0.213, and at 0.5 M CeO₂, it rises to 0.574. The efficiency of CeO₂ as a radiation attenuating material is demonstrated by the fact that an increase in CeO₂ concentration raises the absorption capacity at

Table 1. Experimental values of LAC and MAC for cerium dioxide thin films at different molarity rate for the different K alpha energies.

	Samples	Energy (keV)				
		14.957	22.162	32.191	41.529	59.543
LAC (cm ⁻¹)	0.05 M CeO ₂	1.627 ± 0.131	0.622 ± 0.050	0.226 ± 0.018	0.070 ± 0.006	0.069 ± 0.006
	0.1 M CeO ₂	1.635 ± 0.131	0.625 ± 0.050	0.261 ± 0.021	0.080 ± 0.006	0.078 ± 0.006
	0.2 M CeO ₂	1.637 ± 0.134	0.675 ± 0.054	0.267 ± 0.021	0.087 ± 0.007	0.086 ± 0.007
	0.3 M CeO ₂	1.677 ± 0.137	0.688 ± 0.055	0.282 ± 0.023	0.090 ± 0.007	0.088 ± 0.007
	0.4 M CeO ₂	1.713 ± 0.140	0.700 ± 0.056	0.308 ± 0.025	0.105 ± 0.008	0.097 ± 0.008
	0.5 M CeO ₂	1.747 ± 0.141	0.720 ± 0.058	0.351 ± 0.028	0.126 ± 0.010	0.099 ± 0.008
MAC (cm ² /g)	0.05 M CeO ₂	10.260 ± 0.821	3.902 ± 0.312	1.417 ± 0.113	0.442 ± 0.035	0.432 ± 0.035
	0.1 M CeO ₂	10.415 ± 0.833	3.974 ± 0.318	1.663 ± 0.133	0.512 ± 0.041	0.493 ± 0.039
	0.2 M CeO ₂	10.699 ± 0.856	4.309 ± 0.345	1.703 ± 0.136	0.555 ± 0.044	0.547 ± 0.044
	0.3 M CeO ₂	10.750 ± 0.860	4.315 ± 0.345	1.770 ± 0.142	0.567 ± 0.045	0.555 ± 0.044
	0.4 M CeO ₂	10.942 ± 0.875	4.385 ± 0.351	1.927 ± 0.154	0.656 ± 0.052	0.609 ± 0.049
	0.5 M CeO ₂	11.200 ± 0.896	4.585 ± 0.367	2.237 ± 0.179	0.801 ± 0.064	0.632 ± 0.051

every energy level [44, 45]. Furthermore, studies have demonstrated that a larger mass attenuation coefficient value is necessary for improved x-ray and gamma-ray shielding efficacy [46].

HVL, MFP and TVL values, which are another important radiation shielding parameters, were obtained for five different radiation energy levels from 14.957 KeV to 59.543 KeV for six different molarities and are presented in table 2. Additionally, graphs of these values are given in figures 7(a)–(c). HVL is defined as the thickness of the material at which the intensity of the incoming radiation is reduced by half and TVL is defined as the thickness of the material at which the intensity of the incoming radiation is reduced by one-tenth. MFP is the average distance that radiation can travel in a substance before any two successive collisions [47]. When figure 7(a) was examined, it was seen that the HVL values increased significantly with increasing energy. For example, for 0.05 M, the HVL value was 0.42 cm at the lowest energy of 14.957 keV, while it reached approximately 10 cm at the highest energy of 59.543 keV. This shows that higher energy radiation can penetrate deeper into the material [48, 49]. In addition, as can be seen from table 2, it was determined that there was a decrease in HVL values with the increase in molarity in each energy. It was observed that with the increase in molarity at 14.957 keV energy, the HVL value decreased from 0.424 cm to 0.394 cm, while at 22.162 keV energy, from 1.11 cm to 0.96 cm, at 32.191 keV energy, from 3.07 cm to 1.97 cm, at 41.539 keV energy, from 9.85 cm to 5.51 cm and finally at 59.543 keV energy it decreased from 10.06 cm to 6.98 cm. These results clearly show that higher concentrations of CeO₂ increase the absorption of radiation [44]. However, the results indicate that high-energy radiation penetrates less dependent on material thickness [50]. Similarly, when the MFP values given in figure 7(b) were examined depending on the radiation energy and CeO₂ molarity, distinct trends affecting the material-radiation interactions were observed. When the radiation energy sent to the material was increased from 14.957 keV to 59.543 keV, it was observed that the MFP values increased steadily from 0.61 cm to 14.51 cm (0.05 M CeO₂) This is attributed to the higher energy radiation traveling longer distances [51]. However, with the increase in CeO₂ molarity, more radiation was absorbed or scattered for all energies, and thus a decrease in MFP values was observed.

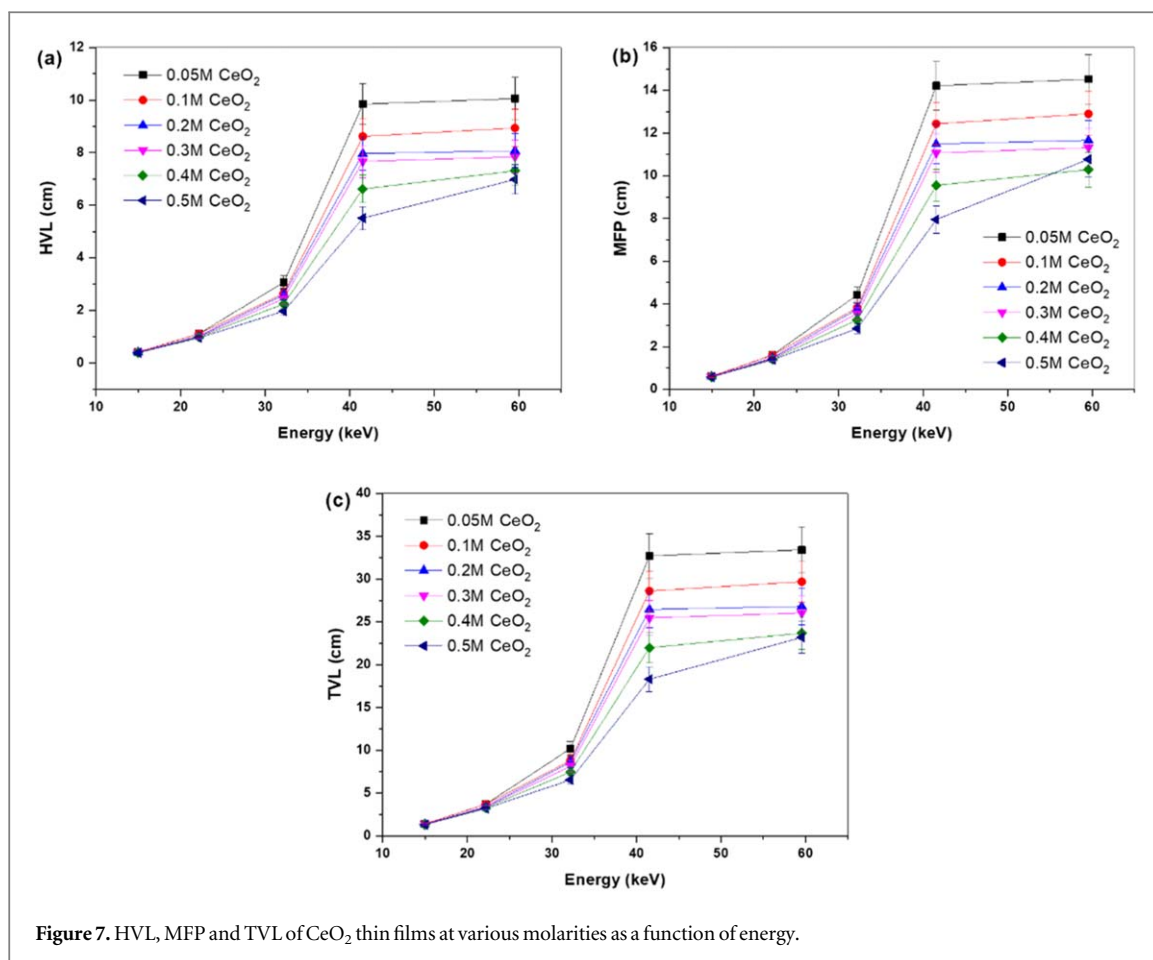
Herein, the lowest MFP values were observed at the low energy level of 14.957 keV and the highest molarity value (0.5 M), revealing the dominance of absorption mechanisms in a dense environment. In contrast, the highest MFP values were obtained in high energy (59.543 keV) and low molarity (0.05 M) samples because radiation-material interactions were determined to be at a minimum level in these conditions. In figure 7(c), TVL values are presented depending on the radiation energy and CeO₂ molarity. The increase in the energy level from 14.957 keV to 59.543 keV decreased the possibility of radiation interacting with the material, which caused the TVL values to increase. This shows that high energy photons travel longer distances in the material without being weakened. On the other hand, the increase in CeO₂ molarity increased the density of the material, causing the radiation to interact more frequently and decreasing the TVL values. The lowest TVL values indicate that low energy radiation (14.957 keV) is rapidly absorbed in a high density (0.5 M) medium and an effective attenuation occurs. In contrast, a high energy (59.543 keV) and low density (0.05 M) medium produced the highest TVL values of 33.423 cm, allowing the radiation to pass through the material with minimal interaction.

Calculations and measurements were carried out by considering the margins of error given in table 3. From the results, it can be concluded that CeO₂ thin films at higher molarities provide better radiation absorption properties and thus may be potential candidates for radiation shielding applications.

Table 2. Experimental values of MFP, HVL, TVL for cerium dioxide thin films at different molarity rate for different K alpha energies.

Samples	Energy (keV)								
	14.957			22.162			32.191		
	MFP	HVL	TVL	MFP	HVL	TVL	MFP	HVL	TVL
0.05 M CeO ₂	0.612 ± 0.049	0.424 ± 0.034	1.408 ± 0.113	1.608 ± 0.129	1.114 ± 0.089	3.702 ± 0.296	4.429 ± 0.354	3.070 ± 0.246	10.197 ± 0.816
0.1 M CeO ₂	0.611 ± 0.049	0.423 ± 0.034	1.407 ± 0.113	1.601 ± 0.128	1.110 ± 0.089	3.687 ± 0.295	3.825 ± 0.306	2.651 ± 0.212	8.808 ± 0.705
0.2 M CeO ₂	0.596 ± 0.048	0.413 ± 0.033	1.373 ± 0.110	1.481 ± 0.118	1.026 ± 0.082	3.409 ± 0.273	3.746 ± 0.300	2.596 ± 0.208	8.625 ± 0.690
0.3 M CeO ₂	0.584 ± 0.047	0.405 ± 0.032	1.344 ± 0.108	1.454 ± 0.116	1.008 ± 0.081	3.348 ± 0.268	3.545 ± 0.284	2.457 ± 0.197	8.163 ± 0.653
0.4 M CeO ₂	0.572 ± 0.046	0.397 ± 0.032	1.318 ± 0.105	1.428 ± 0.114	0.990 ± 0.079	3.289 ± 0.263	3.250 ± 0.260	2.253 ± 0.180	7.484 ± 0.599
0.5 M CeO ₂	0.569 ± 0.045	0.394 ± 0.032	1.309 ± 0.105	1.389 ± 0.111	0.963 ± 0.077	3.198 ± 0.256	2.847 ± 0.228	1.973 ± 0.158	6.554 ± 0.524

Samples	Energy (keV)					
	41.529			59.543		
	MFP	HVL	TVL	MFP	HVL	TVL
0.05 M CeO ₂	14.211 ± 1.137	9.850 ± 0.788	32.721 ± 2.618	14.515 ± 1.161	10.061 ± 0.805	33.423 ± 2.674
0.1 M CeO ₂	12.432 ± 0.995	8.617 ± 0.689	28.626 ± 2.290	12.902 ± 1.032	8.943 ± 0.715	29.709 ± 2.377
0.2 M CeO ₂	11.488 ± 0.919	7.963 ± 0.637	26.453 ± 2.116	11.654 ± 0.932	8.078 ± 0.646	26.835 ± 2.147
0.3 M CeO ₂	11.065 ± 0.885	7.669 ± 0.614	25.477 ± 2.038	11.311 ± 0.905	7.840 ± 0.627	26.045 ± 2.084
0.4 M CeO ₂	9.552 ± 0.764	6.621 ± 0.530	21.994 ± 1.760	10.289 ± 0.823	7.132 ± 0.571	23.692 ± 1.895
0.5 M CeO ₂	7.953 ± 0.636	5.512 ± 0.441	18.312 ± 1.465	10.077 ± 0.806	6.985 ± 0.559	23.204 ± 1.856

**Table 3.** Margins of error in calculating parameters.

Nature of uncertainty	Uncertainty [%]
The original and attenuated photon intensity	≤5
Photon detection statistics	≤3
Specimen thickness	≤2

4. Conclusions

In this study, the structural, morphological and radiation shielding properties of CeO₂ thin films prepared at different molarities were investigated. XRD pattern analyses showed that the crystal structure of the films was consistent with the literature and that the crystal sizes increased with the increase in molarity. FESEM images revealed that the films were spread homogeneously on the surface and had a granular structure. The thickness of the thin films increased with molarity and this increase contributed positively to the radiation shielding parameters of the films. In particular, it was observed that LAC and MAC values increased with molarity, while HVL, MFP and TVL values decreased. This has proved that CeO₂ thin films can effectively absorb low-energy radiation and exhibit better radiation shielding performance at high molarities. As a result, CeO₂ thin films can be considered as a potential material for radiation protection applications.

CRedit authorship contribution statement

Imran Kanmaz: Conceptualization, Methodology, Validation, Investigation, Visualization, and Writing–Review, Editing and Finalizing. **Oğuzkağan Köksal:** Calculations. **Erhan Cengiz:** Supervision. **Gökhan Apaydın:** Supervision. **Murat Tomakin:** Supervision. All authors discussed experiments and results, have reviewed and edited the manuscript. All authors have given approval for the final version of the manuscript.

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Data availability statement

All data that support the findings of this study are included within the article (and any supplementary files).

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