Short Communication

Optical Investigations of MgO Quantum Dots under Pressure Effect at Near-infrared Region

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Abstract

MgO quantum dots (MgO QDs) are prepared using chemical bath deposition technique. The prepared MgO QDs are characterized by UV-visible (UV-vis) spectrophotometry, Fourier Transformation-Infrared (FTIR) and X-ray diffraction (XRD). The crystallite size goes up as pressure increases and corresponding to transmittance. The structural properties of XRD are elaborated in terms of different pressures, these properties of MgO QDs at different pressures from 50 to 300mbar have been investigated. The crystallite size is varied from 4.3nm to 7.2nm to 12.5nm for 200mbar, 250mbar and 300mbar, respectively. The XRD peaks are displayed at 2θ=38.1º, 42.8º, 31.3º and 34.5º that correspond to (111), (002), (100) and (002) planes, respectively. Also, the optical investigations of UV-vis and FTIR are detailed to understand the behavior of MgO QDs for different pressures. The transmittance, 200-1,200nm is in terms of wavelength at different pressures; 50mbar, 100mbar, 150mbar, 200mbar, 250mbar and 300mbar, while the high transmittance for more than 80% is exhibited at 200-300mbar. To investigate bandgap, it depends on the transmittance. At 200mbar and 423K, the bandgap’s nature has direct inter-band transition. The estimated bandgap is 5.01eV. The bandgap variation is attributed to non-stoichiometry of deposited layer. The FTIR spectra for MgO QDs at different pressures; 50-300mbar are researched to recognize many absorption peaks at different lengths indicating to varied phenomena of un-oxide sized, C-O stretching mode or chemical bonding of Mg-O production.

Keywords: MgO, quantum dots, X-ray diffraction, UV

1 INTRODUCTION

The exploration of nanoscale materials has become a focal point of scientific inquiry, driven by the desire to understand their unique properties and potential applications. Among these nanostructures, quantum dots (QDs) have garnered significant attention for their distinct quantum confinement effects, which give rise to size-dependent electronic and optical properties. In this context, magnesium oxide quantum dots (MgO QDs) stand out as a promising subject for investigation due to their wide bandgap and remarkable stability. The optical properties of MgO QDs have been extensively studied, revealing intriguing phenomena such as quantum confinement effects, size-dependent energy bandgaps and enhanced photoluminescence. These characteristics make MgO QDs highly attractive for various technological applications, including sensors, light-emitting devices and solar cells. However, despite the considerable advancements in understanding their behavior under ambient conditions, the response of MgO QDs to external stimuli remains a relatively unexplored frontier. One crucial external factor is pressure that can significantly influence the properties of nanomaterials. The application of pressure induces a change in the interatomic distances, alters the electronic structure,
which can lead to modifications in the optical response of materials. Investigated MgO QDs under pressure is particularly relevant for expanding our comprehension of their fundamental behavior and unlocking potential applications in high-pressure environments.

Recently, Krishna et al. have investigated the electronic and transport characteristics of selective edge hydrogenated MgO nanoribbons. The transport calculations have revealed that the bare Mg-edge (B-MgO-H) nanoribbon exhibits ohmic current behavior, while the other nanoribbons exhibit negative differential resistance (NDR) behavior. The heterostructure-magnesium oxide-boron (H-MgO-B) device exhibits a significant NDR behavior with a significantly high peak-to-valley current ratio (PVC) of about 3.07×10^5. They have suggested that magnesium oxide nanorods (MgONRs) can be used to design nano-interconnects and NDR based nano-electronic device design by tailoring the edge hydrogenation. While, Hazanka and Kalita have researched the effects of homo- and hetero-type co-doping of B, C and N on the structural and electronic properties of MgO monolayers (MLs). Most of co-doping combinations introduce magnetism to the pristine MgO ML. Hetero-type monolayers prefer to be in magnetic states than homo-type MLs to exhibit higher magnetic moments, with a maximum value of 4μB for B-N co-doping. The overall tuning of the electronic properties renders the co-doped MgO monolayers effective for applications in spintronics and optoelectronics. And, Al-Douri et al. have used the full potential-linearized augmented plane wave (FP-LAPW) method to calculate the indirect energy gap (Γ-X) using density functional theory (DFT). The Engel-Vosko generalized gradient approximation (EV-GGA) and modified Becke Johnson (mBJ) formalisms are utilized to optimize the corresponding potential for energetic transition and optical properties calculations of chalcogenides alloys as a function of quantum dot diameter; it has been used to test the validity of quantum dot potential model. They have investigated the refractive index and optical dielectric constant to explore best applications for solar cells.

Near-infrared (NIR) spectroscopy emerges as a powerful tool for probing the electronic and vibrational transitions of materials in the infrared range. The utilization of NIR spectroscopy in studying MgO QDs under pressure provides a unique opportunity to gain insights into their response to external stimuli at the molecular and electronic levels. By focusing on the near-infrared region, the researchers can explore subtle changes in the electronic structure and vibrational modes of MgO QDs, offering a comprehensive understanding of their behavior under pressure. The study of MgO QDs under pressure is not only significant from a fundamental perspective but also holds practical implications. Many applications in the field of nanoelectronics and optoelectronics, involve materials subjected to varying pressure conditions. Understanding how MgO QDs respond to pressure can contribute to the design and optimization of devices that operate in high-pressure environments. In this context, this research aims to systematically investigate the optical properties of MgO QDs under the pressure effect of using near-infrared spectroscopy. By applying pressure as a controlled parameter, we seek to unravel the intricate interplay between quantum confinement effects and external pressure, shedding light on the underlying mechanisms governing the behavior of MgO QDs. The outcomes of this work have the potential to extend the current understanding of nanoscale materials and pave the way for the development of advanced technologies that leverage the unique properties of MgO QDs under diverse environmental conditions. In summary, the exploration of MgO QDs under pressure effect using near-infrared spectroscopy represents a compelling avenue for advancing our understanding of nanomaterials and harnessing their potential in real-world applications. The intricate interplay between quantum confinement effects and external pressure offers a rich scientific landscape to explore, with implications ranging from fundamental condensed matter physics to applied nanotechnology.

2 EXPERIMENTAL

The synthesis of MgO QDs can be achieved through chemical bath deposition technique, offering different control over crystallite size and properties. For a simple chemical precipitation method for synthesizing MgO QDs, it is necessary to bring magnesium chloride, magnesium nitrate, NaOH, deionized water and ethanol. The followed experimental procedure is:

1. Preparation of Magnesium Precursor Solution: Dissolve the magnesium precursor in deionized water to create a clear solution. Stir the solution thoroughly to ensure complete dissolution.

2. Addition of Sodium Hydroxide: Slowly add a NaOH solution to the magnesium precursor solution, while stirring continuously. This will result in the formation of a white precipitate of Mg(OH)₂.

3. Formation of MgO Nanoparticles: Continue stirring the mixture for a specified period to allow the magnesium hydroxide to form. Adjust the reaction time based on the desired size of MgO nanoparticles.

4. Washing and Drying: (i) Collect the formed Mg(OH)₂ precipitate through filtration or centrifugation. (ii) Wash the precipitate with deionized water to remove impurities and excess reactants. (ii) Optionally, repeat the washing step with ethanol to enhance purity. (iv) Dry the resulting Mg(OH)₂ material, typically using an oven or other appropriate drying techniques.

5. Thermal Decomposition: (i) Heat the dried Mg(OH)₂ precursor at an elevated temperature (e.g., in a furnace) to thermally decompose it into MgO nanoparticles. (ii) The decomposition temperature can influence the final properties of the MgO nanomaterial.

6. Characterization: Characterize the synthesized MgO nanomaterial using UV-visible (Jobin Yvon model HR 800 UV system, Kyoto, Japan), Fourier transformation-
infrared (FTIR) (SHIMADZU (Shimadzu Corporation), China) and X-ray diffraction (XRD) (Philips PW 1710 X-ray diffractometer, USA) techniques.

In the realm of nanotechnology, the synthesis of MgO QDs under the influence of pressure represents an intriguing avenue for exploration, particularly in the context of NIR spectroscopy. Applying pressure during synthesis introduces a dynamic dimension to the process, altering the material’s structural and optical properties. The experiment involves subjecting magnesium precursors to varying pressure conditions, thereby manipulating the quantum dot formation. Under high pressure, the size of MgO QDs is expected to undergo notable changes, influencing their optical characteristics. The near-infrared spectral region is particularly relevant due to its significance in biomedical and telecommunications applications. The pressure-induced alterations in quantum dot size and structure may lead to tunable NIR absorption and emission properties, essential for optimizing their performance in these applications. This experimental synthesis under pressure effect not only expands our understanding of QDs formation but also holds promise for tailoring MgO QDs with enhanced NIR features, paving the way for innovative technological applications.

3 RESULTS AND DISCUSSION

3.1 Structural Properties

The crystallinity of MgO QDs is analyzed using XRD. This technique is employed to give an indication about the grain size of the prepared material. Figure 1A illustrates the XRD results for MgO QDs at 50mbar and 423K. In this diffraction pattern, the peaks at 2θ=37.4º and 42.9º are corresponding to (111) and (002) planes. This indicates the formation of oxide molecules for used material in their corresponding films. Also, there are a small amount of unoxidized atoms mixed with oxide result in the formation of Mg for (100) and (002) planes.

The XRD patterns at oxidation pressures, 100mbar to 150mbar are shown in Figure 1B and C at 423K. From the first growth conditions, it is recognized that the peaks...
are appeared at $2\theta=38.1^\circ$, $42.8^\circ$, $31.3^\circ$ and $34.5^\circ$ that correspond to (111), (002), (100) and (002) planes, respectively. Figure 1D shows the diffraction pattern at 200mbar, where the structure has clearly improved a significant increasing in peak intensity at (111) plane and reduction in peak intensity at other planes. This indicates the formation of nearly stoichiometry MgO.

The intensities of these peaks are reduced as pressure increases to 250mbar as shown in Figure 1E and 1F.

The results below ensure that the optimum value of pressure is 200mbar. The deviation of XRD peak with respect to standard American Society for Testing and Materials (ASTM) data is attributed to mechanical micro stress produced by different recourses like impurities, defects and vacancies reside in the structure. The data at higher pressure, 300mbar and 423K are presented in Table 1. So, the peaks are appeared at $2\theta$ are 38.02$^\circ$ and 41.9$^\circ$, which correspond to (111) and (002) planes. The MgO (002) is a weak, while (111) reflection is a stronger \[16,17\].

We can deduce that the increasing of pressure improves the crystal quality and can introduce two completely processes; improving stoichiometry and decreasing of kinetic energy for the reactive particles in the catholic plasma due to high pressure. The structural properties for each diffraction peak are given in Table 1. This gives different relative intensities, Bragg’s angle and miller indices. While, Table 2 presents full width at half maximum (FWHM) of XRD that corresponding to crystallite size at different pressures. It appears an enhancement with increasing pressure up to 200mbar. Here, we can recognize that the reduction in FWHM will reflect the increasing in crystallite size using Scherrer’s formula\[11\].

\[D = \frac{K\lambda}{\beta \cos \theta}\] (1)

where $K$ is geometric factor (0.9), $\lambda$ is XRD wavelength (1.54056Å for Cu-Ka), $\beta$ is FWHM of diffraction peak and $\theta$ is Bragg’s diffraction angle.

### 3.2 Optical Properties

The pressure effect on optical properties of MgO QDs is investigated extensively. The transmittance and bandgap are measured as plotted in Figure 2. The transmittance spectra are recorded in the ultraviolet, visible and NIR ranges at different pressures. In general, it has been found that the transmittance is improved when pressure increases. From Figure 2A, the optical transmittance,
200-1,200nm as a function of wavelength at different pressures is shown, the high transmittance, >80%, is exhibited at 200-300mbar, however, the reduction in transmittance at NIR wavelengths is related to reflection of their photons due to interaction with plasma in conduction band [18].

The bandgap depends on the transmittance. The plots of (α\(h\nu\))^2 against \(h\nu\) at 200mbar and 423K is illustrated in Figure 2B. The nature of plots suggests a direct inter-band transition. The estimated bandgap is 5.01eV. It can state that a reduced bandgap is attributed to varied extent of non-stoichiometry of the deposited layer [19].

The transmittance as a function of deposition position upon the substrate surface is illustrated in Figure 3. It is known that the transmittance of incident 632.8nm wavelength decreases to its minimum value whenever approaching to the center, where the lowest transmittance can be achieved. This relates to the thickness, which is higher at the center and reduces as we depart from the radial direction. This coincides with the theoretical concept that imposes the deposition or evaporating an atom from the target with a solid angle. This result ensures the difference in the transmitted power that can be neglected for a limited area, 1cm², which found to have a uniform transmittance that attributed to the crystalline nature throughout the coated area, which is obtained a lattice arrangement for resulting in the better optical properties [20].

The Fourier Transformation-Infrared spectroscopic results give information about phase composition and the way in which oxygen is bound to metal ions. The following show FTIR spectra for MgO QDs at different pressures. At different pressures; 50-300mbar, we can see the change in the composition. Figure 4A gives FTIR result at 50mbar, we can recognize the absorption peak at 1,620.9, 786.9, 725.18cm\(^{-1}\) that is related to Mg atoms, beside 694.33, 648.04cm\(^{-1}\) absorption spectra are connected to MgO QDs. An increasing in MgO QDs formation could be recognize by increasing of pressure to 100mbar as shown in Figure 4B where 694.33, 609.46, 408.88cm\(^{-1}\) absorption peak could be found. In addition, the presence of 987.49, 918.05, 817.7cm\(^{-1}\) peak indicates to un-oxide sized. The peak at 1,465.8, 1,049.2 and 871.76cm\(^{-1}\) at low pressure in 50-100mbar is related to C-O stretching mode, which may be attributed to the fact that Mg is fired rather than react with O\(_2\) atoms. At high pressure, 150mbar (Figure 4C), it is recognized that the peak is eliminated to one peak at 918.05cm\(^{-1}\), while higher number of MgO QDs is formed, this could be obtained at 648.04, 466.03, 408.88cm\(^{-1}\) at 200mbar (Figure 4D), the absorption spectra is clearly disappeared while it is found at 655.75, 570.89, 455.17, 408.88cm\(^{-1}\). For MgO QDs, the kinetic energy of Mg atoms decreases through collision with oxygen, so the chemical bonding like (Mg-O) is produced. By increasing the pressure to 250 and 300mbar, Figure 4E and 4F shows that the peaks at 655.75, 586.32, 501.46, 408.88cm\(^{-1}\) are related to MgO QDs. In all above result, it is recognized that the peaks at 599.03, 879.48, 840.91, 1,396.37, 1,342.36, 1,072.19, 1,612.38, 1,550.66, 1,296.08, 1,064.63cm\(^{-1}\) are related to glass substrate [21-31].

4 CONCLUSIONS
The chemical bath deposition technique was proved to be an effective method for synthesizing MgO QDs. The pressure effect has been employed to investigate different structural and optical studies of MgO QDs. The
Scherrer’s formula has been utilized to research structural properties that proved the MgO QDs has the highest peak (111). Different pressures; 50mbar, 100mbar, 150mbar, 200mbar, 250mbar and 300mbar were applied. The pressure, 200mbar has presented as evidence to indicate the ideal intensity for investigating referring to 4.3nm crystallite size. Many peaks corresponding to many planes were displayed in XRD pattern. All of these were supported by studying the optical properties of transmittance. The energy bandgap and FTIR were to confirm the fact that 5.01eV was close to NIR, the nature of bandgap was direct inter-band transition and attributed to non-stoichiometry of deposited layer, and the correlation between transmittance and pressure was directly. In addition, the transmittance was more than 80% at 200-300mbar. The FTIR spectra for MgO QDs at different pressures; 50-300mbar were recognized many phenomena of un-oxide sized, C-O stretching mode or chemical bonding of Mg-O production.

**Figure 4. FTIR spectrum of MgO QDs.** A: FTIR spectrum of MgO QDs at 50mbar; B: FTIR spectrum of MgO QDs 100mbar; C: FTIR spectrum of MgO QDs 150mbar; D: FTIR spectrum of MgO QDs 200mbar; E: FTIR spectrum of MgO QDs 250mbar; F: FTIR spectrum of MgO QDs 300mbar.
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Conflicts of Interest
The authors declared no conflict of interest.

Author Contribution
Al-Douri Y wrote, edited, and reviewed the original draft, while also copyrighting the figures and tables. Ameri M organized and validated the manuscript, and Bouhemadou A edited and reviewed the manuscript.

Abbreviation List
FTIR, Fourier transformation-infrared
FWHM, Full width at half maximum
MgO QDs, MgO quantum dots
MLs, Monolayers
NDR, Negative differential resistance
NIR, Near-infrared
QDs, Quantum dots
UV-vis, UV-visible
XRD, X-ray diffraction

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