Research Article

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Photocatalytic Degradation of Transition Metal (Ni) Doped ZnS Nanoparticles Synthesized via Simple Solvothermal Route

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ABSTRACT

Ni doped ZnS nanoparticles were obtained using the simple Solvothermal Microwave Irradiation (SMI) method in this research. This technique is cost-effective and results in a high level of uniformity in particle size. The sample's structural characteristics were examined utilizing XRD Technique, FESEM image, Elemental analysis of the as prepared sample obtained from EDX spectrum and their optical characteristics analysed by using UV-Visible absorption study. The XRD pattern of Ni doped ZnS nanoparticles were to obtain their crystallite size (D), lattice constant (a), and volume of the unit cell (V). As the concentration of Ni dopant increased (0, 2.5, 5.0) M %, the values of the lattice constant decreased, leading to an overall decrease in the volume of the unit cell. FESEM images reveals the sample have uniform surface morphology and EDX analysis give evidence for element Zn, Ni, S presents in the sample. Ni doped ZnS nanoparticle dimension strongly influences their optical characteristics. The optical bandgap, as obtained from UV-Vis measurements, ranges from 3.54 eV to 3.50 eV with doping concentrations varying from 0 M% to 5.0 M% .This adjustment in optical properties suggests that Ni-doped ZnS nanoparticles have diverse applications in optoelectronics, sensors, UV detectors, and as an efficient photocatalyst for degrading pollutants in water. The efficiency of the degradation of Methylene Blue dye by Ni doped ZnS nanoparticles was found to be 75.19%.

Keywords: Ni doped nanoparticles; SMI technique; XRD analysis; UV-Visible analysis; Photocatalyst.

1. INTRODUCTION

Now a day's searching a new material have multidisciplinary properties. It has the quality to overcome all our environmental and human needs. The scientist's society only will find new multidisciplinary functional materials to resolve the pollutants in water and environments. Group II-IV chemical compounds give large expectation due to their unique qualities such as morphological and optical properties (Krishnan *et al*.2019; Rajabi and Farsi, 2015). The broad band gap characteristics of the non-toxic ZnS nanomaterial (3.0-3.77 eV) are essential for optoelectronic applications. These attributes include optical, electrical, and photocatalytic qualities that are essential for the creation of solar cells and other optoelectronic devices.

Various physical and chemical methods, such as hydrothermal processes (Wu *et al.* 2017), Sonochemical (Amiri *et al*. 2014), Chemical bath deposition (Zeinand Alghoraibi, 2019), Solid State method (Farooqi and Srivastava, 2014), Coprecipitation method (Srivastava *et al*. 2013), Microwave assisted irradiation method (La *et al.* 2013), Solvothermal Microwave Irradiation (SMI) technique (Gowdhaman *et al.* 2021) etc were employed to obtained ZnS nanoparticles. The SMI technique is considered one of the most effective ways to produce ZnS nanomaterial because the particles are minimized dimension, have a consistent shape, and the method guarantees safety and high purity (Gowdhaman *et al.* 2021). The customization of nonmaterial properties for device use through the addition of appropriate materials is a well-established practice. When semiconductor nanoparticles are doped with transition elements such as Cd, Ag, Cu, Ni, Co and Mn, new possibilities arise due to entire characteristics change thehost materials ZnS (Ramasamy *et al.* 2012). Scholars have endeavoured in diverse ways to incorporate metal ions into ZnS nanoparticles.

Through the process of doping, significant enhancements have been made to the fundamental physical characteristics of ZnS, particularly for the purposes of optoelectronic device utilization (Xu *et al.* 2018). The inclusion of Cu in ZnS material enables precise band gap manipulation, expanding its potential for use in high-performance optoelectronic device components such as solar cell windows and in the photocatalytic degradation of industrial dyes showed by (Wang *et al.* 2019). Transition metal (Fe) doped ZnS nanoparticles in photocatalytic activity on Turquoise Blue H5G dye is superior to that of pure ZnS reported by

(Suganthi and Pushpanathan, 2019). Controlled precipitation method used to doped Cu (0.1%) with ZnS for study their characteristic change. Cu (0.1%) doped ZnS nanoparticles reported optical bandgap value E_g changes from 4.41-4.47 eV. The average of particles size of the as synthesized sample was found to be about 2.0–10.0 nm reported by (Pouretedal and Keshavarz, 2010).

With all of the aforementioned qualities of ZnS nanoparticles, making them a desirable option for multiple applications in electronics. In ZnS, Ni impurities are seen as ideal acceptors, adjusting its optical characteristics. When Ni ions occupy the Zn lattice sites, they act as sites for capturing charge carriers, which prevents the recombination of charge carriers and improves its efficiency for photocatalytic activity (Kaur *et al*. 2015). (Jothibas *et al*. 2018) have used solid state method to prepared doped Ni (0.5-2.0%) with ZnS for tuning the bandgap and the crystallite size also increased with increase dopant. The distance between valance and conduction value (E_g) changed from 3.58 eV to 3.97 eV.

To prepare of pure and (Ni) (0 M%, 2.5 M%, 5.0 M%) doped ZnS nanoparticles was carried out using the simple solvothermal Microwave Irradiation (SMI) method in this study. The research involved a comprehensive examination of the structural, morphological, and optical properties using XRD, FESEM, EDX, HRTEM, and UV visible spectroscopy techniques.

2. EXPERIMENTAL METHODS

For pure and different concentration (2.5 M% and 5.0 M %) of Ni doped ZnS nanoparticles, the reactents Zn(II) acetate dihydrate ($Zn(CH_3COO)_2{\cdot}2H_2O$), Nickel(II) acetate tetrahydrate ($Ni(OCOCH_3)_2{\cdot}4H_2O$), Thiourea ($NH_2CSNH_2$), and ethylene glycol ($C2H_6O_2$) were purchased from Sigma-Aldrich. The cost-effective simple solvothermal microwave irradiation method (SMI) is used to synthesizing the above nanoparticles (Saravanan *et al*. 2012). In order to synthesise 2.5 M% of Ni-doped ZnS nanoparticles, a solution was prepared by dissolving 0.1M of Zn (II) acetate dihydrate and 0.025 M of Nickel (II) acetate tetrahydrate in 50 ml of ethylene glycol. Subsequently, a solution of 0.3 M of thiourea in 50ml ethylene glycol was introduced gradually into the previously described mixture, with continuous stirring. To get a homogeneous solution, the mixture was agitated vigorously for one hour. A double-walled microwave-safe bowl was employed to contain the resultant homogenous solution, approximately 100 ml in volume. The solution was heated in a domestic microwave oven (Model No. IFB 23BC4 23 L) until a colloidal precipitate was obtained. The resulting colloidal precipitate was cooled to ambient temperature and purified with double-distilled water and acetone to remove any remaining contaminants and unreacted starting materials. The same method was used for the synthesis of 5.0 M% Ni-doped ZnS nanoparticles. Subsequently, the gathered nanoparticles were employed for further characterization. To synthesize pure ZnS nanoparticles, a solution was prepared by dissolving 0.1 M of Zn (II) acetate dihydrate in 50 ml of ethylene glycol. A 0.3 M solution of thiourea in 50 ml of ethylene glycol was then added gradually to the aforementioned mixture while maintaining continuous stirring. To achieve a homogeneous solution, the mixture was vigorously agitated for one hour, and the previously described process was employed to obtain pure ZnS nanoparticles.

3. PHOTODEGRADATION OF METHYLENE BLUE (MB) DYE

The synthesized Ni-doped ZnS nanoparticles were evaluated for their photocatalytic activity in degrading methylene blue (MB) dye in aqueous solutions. A stock solution of MB was prepared at a concentration of 10 mg/L, and 30 mg of the as-prepared Ni-doped ZnS nanoparticles was added to 100 mL of this solution. The mixture was stirred using a magnetic stirrer for one hour to ensure uniform dispersion of the nanoparticles and enhance their interaction with the dye molecules. Following this, the combined solution was exposed to ultraviolet (UV) radiation, which excited the electrons in the nanoparticles, generating reactive species capable of degrading the MB dye. Samples were extracted at 30-minute intervals over a total period of 120 minutes and analyzed using a UV-Vis spectrophotometer to measure the absorbance at the dye's characteristic wavelength.

4. RESULT AND DISCUSSION

4.1 Structural Analysis

Figure 1 presents the X-ray diffraction (XRD) patterns of Ni-doped ZnS nanoparticles at doping concentrations of 0 M%, 2.5 M%, and 5.0 M%. The analysis reveals that the synthesized nanoparticles exhibit a cubic ZnS structure, consistent with the standard JCPDS Card No. 05-0566, as reported by Tiwary *et al.* (2021) and Patel et al. (2018). Importantly, no impurities or secondary phases were observed, confirming the successful incorporation of nickel (Ni) into the ZnS lattice. The structural integrity of the Ni-doped nanoparticles is attributed to the smaller ionic radius of Ni (0.69 Å) compared to that of the zinc (Zn) atom (0.74 Å), facilitating its substitution without distorting the cubic framework.

As the Ni doping concentration is increased from 2.5 M% to 5.0 M%, notable changes in the XRD peaks were observed, including alterations in intensity and broadening, which suggest variations in crystallite size and potential strain within the nanoparticles. The XRD data were employed to derive key structural

parameters such as crystallite size, lattice constant, and unit cell volume. Utilizing Bragg's equation ($n\lambda = 2d \sin\theta$), the interplanar spacing was calculated, while the average crystallite size was estimated through Scherer's formula (Tiwary *et al*.2021; Hussein, 2023).

$$D = K\lambda / \beta\cos\theta \quad \ldots\ldots\ldots\ldots (1)$$

Where, K is a dimensionless shape factor it, has a typical value of about 0.9. The Bragg's diffraction angle is represented by 'θ (degree), while the FWHM of diffraction peaks at 2θ (degree) is denoted by β, and λ represents the wavelength (nm) of the X-ray use.

Table 1. The structural parameters of Ni (0, 2.5, and 5.0) M% doped ZnS nanoparticles

Chemical Composition	Crystallite Size(nm)	Inter planar spacing 'd' Å	Lattice Constant a (Å)	Volume of a Unit cell ('a³' (Å)³)
Pure ZnS	7.01	3.198	5.384	155.72
2.5% Ni doped ZnS	6.32	3.634	5.362	153.99
5.0% Ni doped ZnS	6.09	3.762	5.352	149.72

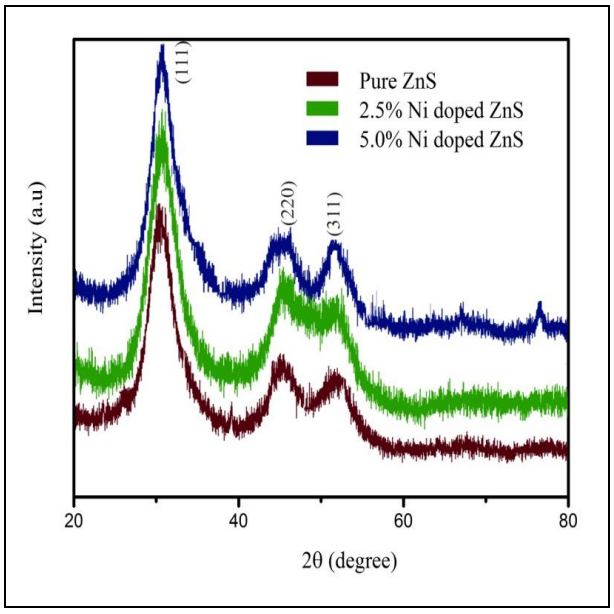

Fig. 1: XRD patterns of Ni (0, 2.5, 5.0) M% doped ZnS nanoparticles

From the obtained structural data, an interesting trend was noted: as the concentration of Ni increased, the lattice constant exhibited a decrease, resulting in a corresponding reduction in the volume of the unit cell. Table 1 summarizes the structural parameters of both pure and Ni-doped ZnS nanoparticles, highlighting the significant effects of doping on their crystallographic characteristics.

The FESEM images (Fig. 2) reveal the structural characteristics of Ni-doped ZnS nanoparticles at doping concentrations of 0 M%, 2.5 M%, and 5.0 M%. Notably, the analysis indicates that while pure ZnS nanoparticles exhibit larger particle sizes, the introduction of nickel as a dopant leads to a reduction in particle size. This size modification can be attributed to the incorporation of Ni ions into the ZnS lattice, which influences the nucleation and growth processes during synthesis. The observed agglomeration of nanoparticles is consistent with findings by Chen *et al.* (2013) and suggests that the dopant may enhance the stability of smaller particle sizes through altered interparticle interactions.

Additionally, the uniform distribution of particle sizes in the Ni-doped ZnS samples can be linked to the solvothermal microwave irradiation synthesis technique employed. This method provides controlled thermal conditions that promote uniform growth and minimizes particle coalescence. The predominance of nearly spherical shapes among the nanoparticles further supports the effectiveness of this synthesis approach, as spherical geometries are typically favoured in processes where energy minimization is a factor. The combination of these factors not only elucidates the role of nickel doping in particle size reduction but also highlights the importance of synthesis techniques in achieving desirable material properties for potential applications in photocatalysis and optoelectronics reported by Kumar *et al*. (2019).

4.2 Elemental Analysis

Figure 3 illustrates the Energy Dispersive X-ray Spectroscopy (EDX) results for pure ZnS nanoparticles alongside ZnS nanoparticles doped with 2.5 M% and 5.0 M% Ni. The EDX analysis confirms the presence of zinc (Zn), nickel (Ni), and sulphur (S) in the samples, demonstrating that the incorporation of nickel does not disrupt the fundamental stoichiometry of the ZnS matrix. Fig. 3 a, b, and c illustrate the EDX (Energy Dispersive X-ray Spectroscopy) findings for pure ZnS nanoparticles as well as ZnS nanoparticles containing 2.5 M% and 5.0 M% Ni doping. Analysis via EDX indicates the existence of Zn, Ni, and S elements while preserving their stoichiometric proportions. This is crucial, as maintaining the stoichiometric proportions ensures that the electronic and optical properties of the nanoparticles remain optimal for applications in photocatalysis and dye-sensitized solar cells. Moreover, the EDX findings reveal insights into the effectiveness of nickel doping in modulating the properties of ZnS nanoparticles. The slight increase in nickel concentration correlates with potential alterations in the band gap and charge carrier dynamics, which could enhance photocatalytic efficiency

(Raza *et al*. 2024). By maintaining the stoichiometric balance between Zn and S while introducing nickel, these nanoparticles are poised to leverage the unique properties of transition metals, such as enhanced charge separation and improved light absorption, thereby broadening their applicability in advanced materials science and energy conversion technologies (Jothibas *et al.* 2018).

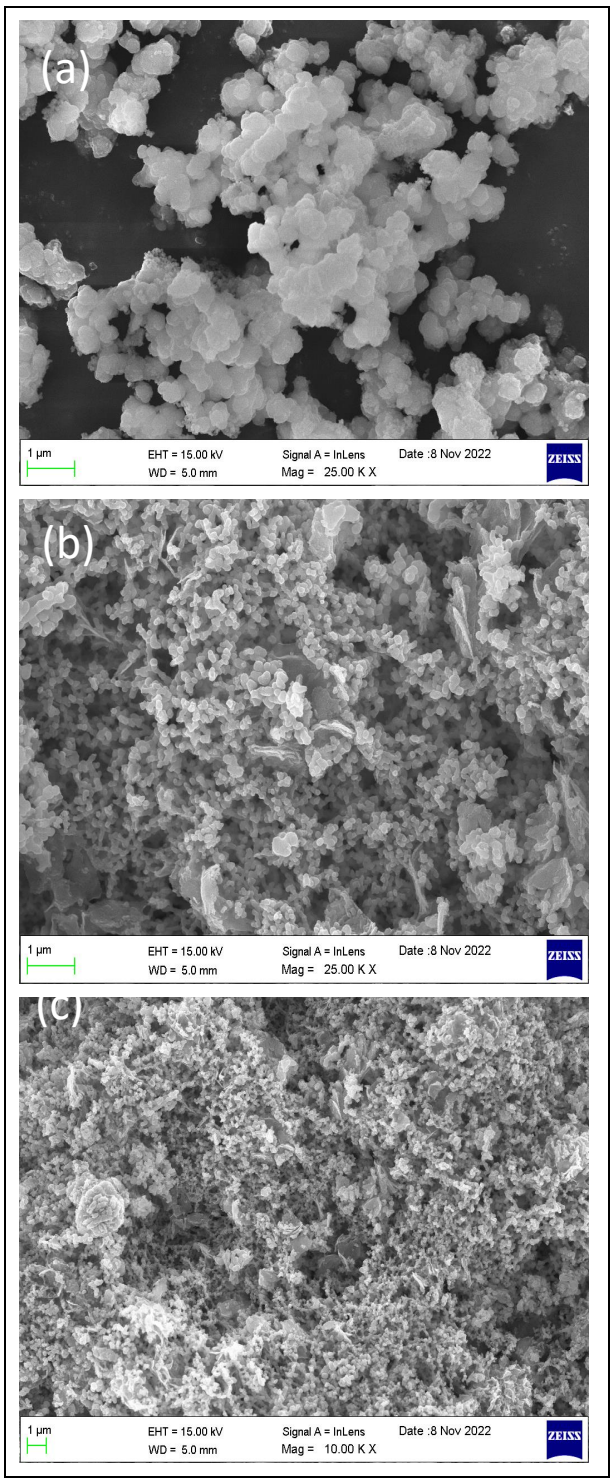

Fig. 2: FESEM images of (a) pure ZnS, (b) 2.5 M% Ni-doped ZnS Nanoparticles, (c) 5.0 M% Ni-doped ZnS nanoparticles

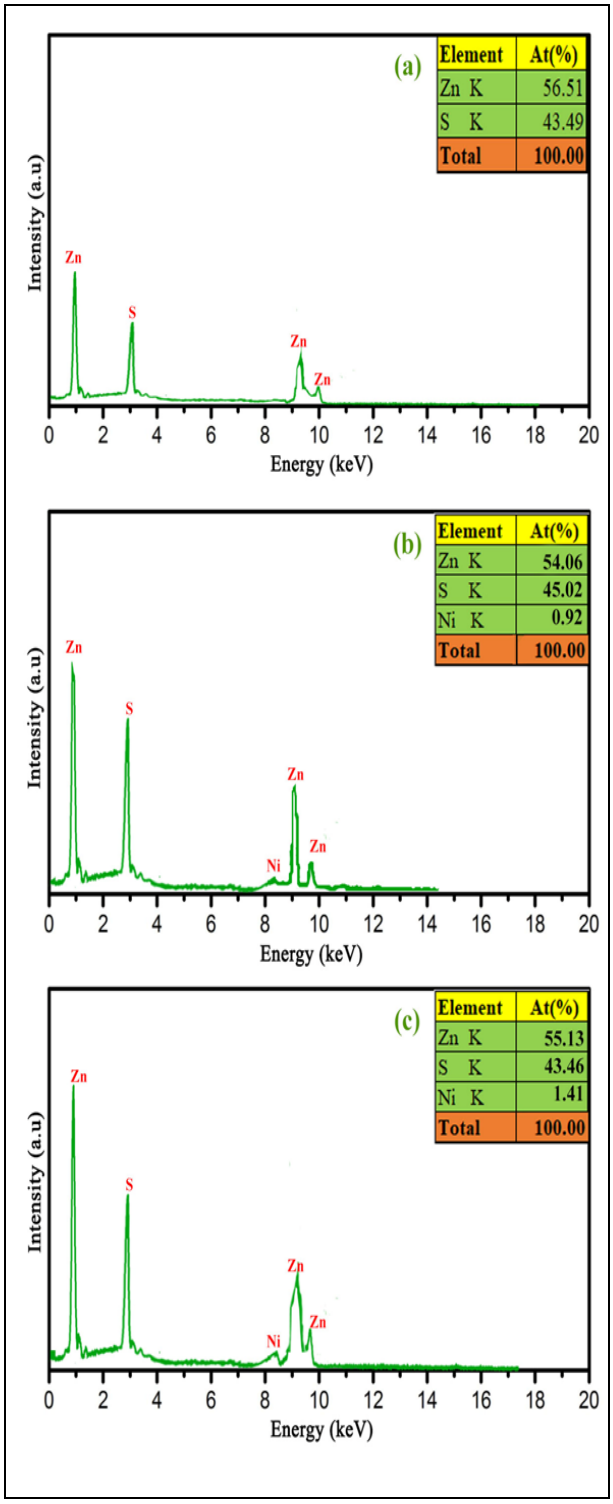

Fig. 3: EDX Spectra of (a) pure ZnS, (b) 2.5 M% Ni-doped ZnS Nanoparticles, (c) 5.0 M% Ni-doped ZnS Nanoparticles

4.3 Optical Studies

The optical properties of Ni-doped ZnS nanoparticles were systematically investigated through UV-visible absorption spectroscopy, with results illustrated in Figure 4(a). Notably, the absorption band

edge was consistently observed between 270 nm and 370 nm across all samples. As the concentration of Ni dopant increased, a red shift in the absorption spectrum was noted, indicating a significant influence of the dopant on the electronic structure of the nanoparticles. This shift can be attributed to the quantum confinement effect, which becomes more pronounced with higher doping levels. The corresponding optical bandgap energies were determined from the energy plots shown in Figure 4(b), revealing values of 3.54 eV, 3.52 eV, and 3.50 eV for 0 M%, 2.5 M%, and 5.0 M% Ni concentrations, respectively. These results suggest that as Ni concentration increases, the optical bandgap decreases, reflecting a higher charge carrier concentration that facilitates enhanced photon absorption.

The decrease in the optical bandgap with increasing Ni dopant levels signifies a transition in the electronic properties of the nanoparticles, likely due to enhanced interaction between the dopant and the ZnS lattice. This increased charge carrier concentration accelerates the absorption of photons, thereby optimizing the photocatalytic potential of the Ni-doped ZnS nanoparticles. The observed modifications in optical characteristics underscore the suitability of these nanoparticles for photocatalytic applications, as they exhibit improved absorption capabilities that are essential for effective energy conversion processes (Joseph *et al*. 2022). Such findings highlight the promising nature of Ni-doped ZnS nanoparticles in various photcatalytic and optoelectronic applications, further enhancing their appeal in advanced material research (Kumar *et al*. 2019).

4.4 Photocatalytic Activity on MB Dye

The performance of photo catalysis is influenced by crystallinity, surface area, and morphology. The MB solution's peak absorbance wavelength was observed at 665 nm (Markovic *et al.* 2015). The process of photocatalysis usually commences with the absorption of photons, leading to generate electron-hole pairs. Afterwards, the charged particles spread to the surface of the sample. On the surface, they react with water, producing reactive peroxide (O_2^-) and hydroxyl radical ($^*OH^-$), which are play role for breaking down dye molecules. The process of photocatalytic activity is as follows.

$$ZnS + hv \longrightarrow e^- + h^+$$

$$e^- + O_2{}^* \longrightarrow O_2^-$$

$$h^+ + OH \longrightarrow {}^*OH$$

$$h^+ + H_2O \longrightarrow H^+ + {}^*OH^-$$

$$^*OH^- + \text{Methylene Blue} \longrightarrow \text{Degradation products}$$

$$^*O^- + \text{Methylene Blue} \longrightarrow \text{Degradation products}$$

The interaction of highly reactive radicals such as *OH and $*O_2^-$ with methylene blue dye is pivotal in the degradation process. These radicals initiate a series of oxidation reactions, breaking down the dye's complex structure into smaller, less harmful molecules. The effectiveness of the photocatalytic activity is quantitatively measured using the percentage of degradation (D), calculated with the formula (Pouretodal *et al*. 2010):

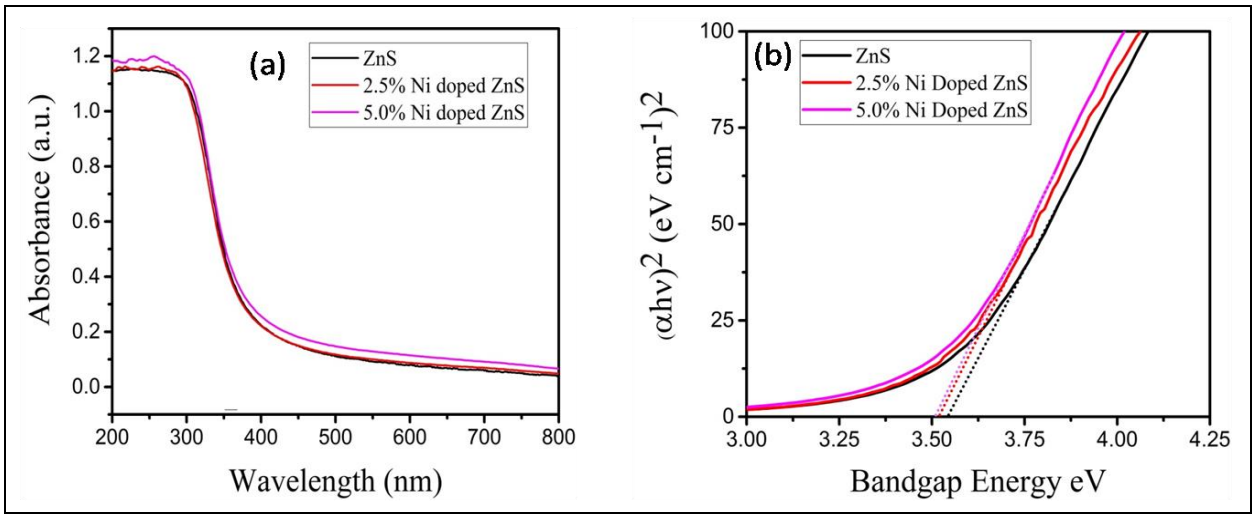

Fig.4: (a) UV-Vis spectra of Ni (0, 2.5, and 5.0) M% doped ZnS nanoparticles (b) Tauc plot for Ni (0, 2.5, and 5.0) M% doped ZnS nanoparticles

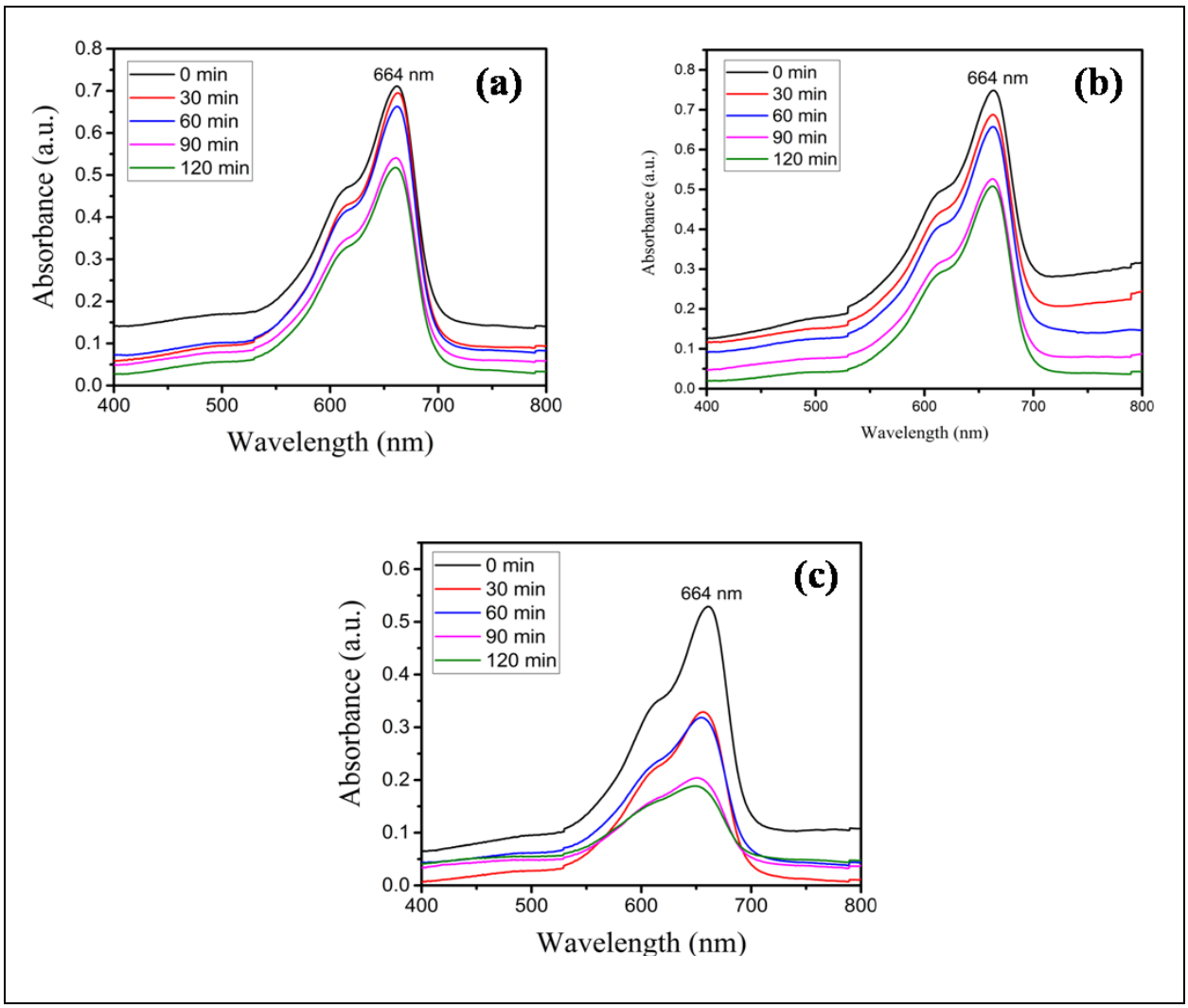

Fig. 5: Photodegradation response of MB in water under UV radiation, (a) Pure ZnS, (b) 2.5 M% Ni, (c) 5.0 M% Ni doped ZnS nanoparticles

$$\% D = [(C_0-C)/C_0]*100 \% \quad \text{--------- (2)}$$

Where C_0 is the initial concentration of the dye and C is the concentration at various time intervals during UV irradiation, up to 120 minutes. This method allows for a clear assessment of the catalyst's efficiency in reducing dye concentration over time.

Experimental results reveal that the degradation efficiencies for pure and Ni-doped ZnS catalysts are 60.2 % for pure ZnS, 64.5% for 2.5 M% Ni-doped ZnS, and an impressive 75.2% for 5.0 M% Ni-doped ZnS nanoparticles. This data indicates a clear trend: as the Ni doping concentration increases, the photocatalytic efficiency improves significantly. The substantial enhancement in degradation efficiency with higher Ni doping can be attributed to several factors. Firstly, the optimal 5.0 M% Ni doping may create more active sites for the adsorption of MB dye, allowing for increased interaction with reactive radicals. Furthermore, the presence of Ni can facilitate better charge separation, reducing recombination rates of electron-hole pairs and thereby increasing the availability of these charge carriers for oxidative reactions (Jothibas *et al.* 2018). This trend underscores the importance of optimizing dopant concentration in enhancing photocatalytic performance, making these Ni-doped ZnS nanostructures particularly effective for environmental remediation applications.

5. CONCLUSION

In this study, we successfully synthesized pure and nickel-doped ZnS nanoparticles at concentrations of 0 M%, 2.5 M%, and 5.0 M% using the simple solvothermal microwave irradiation (SMI) method, which facilitated uniform growth and controlled particle sizes. Characterization techniques, including X-ray diffraction (XRD), field emission scanning electron microscopy (FESEM), energy dispersive X-ray

spectroscopy (EDX), and UV-visible absorption spectroscopy, revealed remarkable structural integrity, with XRD patterns confirming a cubic ZnS structure and the successful incorporation of nickel without impurities. FESEM analysis indicated a reduction in particle size with increased Ni doping, while EDX results confirmed the presence of zinc, nickel, and sulfur, ensuring stoichiometric balance. The optical properties showed a notable decrease in optical bandgap as the Ni concentration increased, which enhanced charge carrier dynamics. This culminated in significant improvements in photocatalytic performance, with the 5.0 M% Ni-doped ZnS nanoparticles achieving a remarkable degradation efficiency of 75.2% for methylene blue dye. This enhancement is attributed to increased formation of reactive species, improved charge separation, and greater active site availability for dye adsorption. These findings highlight the effective modulation of ZnS properties through nickel doping and underscore the potential of Ni-doped ZnS nanoparticles as efficient photocatalysts for environmental remediation, paving the way for further exploration in advanced materials and optoelectronic applications.

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CONFLICT OF INTEREST

The authors declared no conflict of interest in this manuscript regarding publication.

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