



Research Article

Sensitivity of Nanostructured Mn-Doped Cobalt Oxide Films for Gas Sensor Application

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Received: Mar. 19, 2020; **Accepted:** Jun. 20, 2020; **Published:** Jul. 13, 2020

Citation: Ehssan Salah Hassan, Kameran Yasseen Qader, Esraa Hassn Hadi, Sami Salman Chiad, Nadir Fadhil Habubi, and Khalid Haneen Abass, Sensitivity of Nanostructured Mn-Doped Cobalt Oxide Films for Gas Sensor Application. *Nano Biomed. Eng.*, 2020, 12(3): 205-213.

DOI: 10.5101/nbe.v12i3.p205-213.

Abstract

The effect of manganese doped cobalt oxide ($\text{Co}_3\text{O}_4:\text{Mn}$) was investigated by two different ratios (1% and 3%), which were precipitated by spray pyrolysis technique (SPT), and was adopted using a laboratory designed glass atomizer. Glass substrates were used to deposit films on them, heated at a temperature of 420 °C. The structural properties were studied through X-ray diffraction. The results showed that all deposit nanostructured films were polycrystalline and there was a decrease in the preferred reflection intensity along (311) plane resulting in a decrease in the crystallite size. Surface properties were analyzed through atomic force microscopy (AFM), which showed a decrease in the roughness and the particle size growth was a vertical columnar rod. The optical characterization displayed that the transmittance of pure Co_3O_4 nanostructured films was 48% and decreased to 35% for 1% of the Mn concentration, and continued to decrease to 33% with the increase of manganese concentration up to 3%. Optical energy bandgap of pure Co_3O_4 nanostructured films was 1.435 eV and decreased to 1.419 eV for 1% of Mn concentration, and continued to decrease to 1.367 eV with the increase of Mn concentration up to 3%. The highest percentage sensitivity was for the sample doped with 3% Mn, which was about 65%, for NO_2 gas concentration of 600 ppm, at an operating temperature of 200 °C.

Keywords: Manganese, Cobalt oxide, SPT, Crystalite size, Columnar rod, Gas sensor

Introduction

Among the different transition metal oxides (TMO), Co_3O_4 is an imperative p-type semiconductor (coordinate energy bandgap at 1.48 and 2.19 eV) [1, 2]. Generally utilized as an electrochemical gadget [3], heterogeneous impetus [4], gas sensor [5], anode material in lithium batteries [6], and sunlight based vitality safeguard [7].

Recently the consideration regarding transition metal oxides has been taken into account. Metal oxide nanostructures play an essential role in numerous locales of science, physical science, ecological science, and material science [8-10]. These TMO, while falling in the nano estimated administration, are relied upon to have significantly witnessed an increasing interest in the fields; for example, information stockpiling, spintronics biomedicine, and

broadcast communications [11-14]. These features are firmly subject to their dimension and surface properties. The Co_3O_4 shows an ordinary spinel structure, in which Co^{2+} is engaged in tetrahedral destinations and Co^{3+} is engaged in octahedral locales [15]. This is very attractive because of the twists of Co^{2+} particles, with a little commitment from turn circle coupling. Then again, Co^{3+} particles have no perpetual attractive point as a result of the part of 3D levels of the octahedral precious stone field and finish filling of levels [16]. Given the attractive properties of permeable Co_3O_4 small scale 3D squares, the antiferromagnetic Co_3O_4 displays a specific level of ferromagnetism under outside attractive field because of the unpaired particles' closeness to the surface of the nanocrystals and the association of molecules. The unique properties of nanoparticle unequivocally reliant change of shapes and sizes and crystallization, charge heading [17]. Recently, a few procedures have been utilized to enhance the conduct of metal oxides by methods for presenting different dopants [18], shaping nanocomposites with p-n intersection [19], and by tuning the morphology [7], have focused on the electrical and surface properties of nanoparticles to accomplish enhanced reactant, electro-optical, attractive, substance, and physical properties [20]. $\text{Co}_3\text{O}_4:\text{Mn}$ produced by sol-gel spin coat method has polycrystalline nature with degradation in the crystallization with increased manganese concentration, and the output phase is cubic with the appearance of orthorhombic MnO_2 phase. Shan, and Dai [21] explained that the doping of cobalt by magnesium causes a decrease in particle size and most particles had a nano-spherical shape and the size were between 82-244 nm. In this research Mn doped cobalt oxide is prepared to study their physical properties to use these films as a gas sensor [22-24]. The PLAL synthesized MoS_2 NS explained the vast absorbance in Vis region. Photocatalytic activity of $\text{TiO}_2+\text{ZSM}_5-\text{MoS}_2$ nanocomposite was examined with arsenite. $\text{TiO}_2-\text{ZSM}_5-\text{MoS}_2$ nanocomposite displayed nearly 100% arsenic photo-conversion to Usenet. This work aims to consider the effect of deformation on the sensitivity of the deposited films for NO_2 gas.

Experimental

An aqueous solution of 0.1 M of $\text{CoCl}_2 \cdot 6\text{H}_2\text{O}$ supplied from Merck Chemicals (Germany) was used to prepare thin films of Co_3O_4 [25], and 0.1 M of

Manganese chloride $\text{MnCl}_2 \cdot 4\text{H}_2\text{O}$ was used as doping agent with a concentration of 1% and 3% to obtain Mn doped cobalt chloride [26]. A few drops of HCl were added to obtain a homogeneous solution. The spray pyrolysis technique was adopted with an experimental intended glass spray [27]. Glass substrate was used for deposition heated at a temperature of 420 °C. The following optimal status was taken into account, with spray time of 8 s. The spray was sprayed with two limits of 1.5 min to avoid excessive cooling and to prevent cracks. Nitrogen carrier gas was upheld at a pressure of 10^5 Nm^{-2} . The space between the nozzle and the substrate was 28 ± 1 cm, at solution flow rate of 4 mL/min. Film thickness was dignified using gravimetric process and was found to be about 320 nm. Optical transmittance and absorbance employing UV-visible spectrophotometer were verified in the wavelengths range of 300-1100 nm (Shimadzu Company, Japan). High accuracy XRD was utilized to investigate the structural qualifications of thin films employing D8 Advance Bruker system, $\text{CuK}\alpha$ ($\lambda = 0.154056 \text{ \AA}$).

Results and Discussion

XRD spectra of pure and $\text{Mn}:\text{Co}_3\text{O}_4$ nanoparticles are illustrated in Fig. 1(a), (b) and (c), from which the phase and crystalline structure were diagnosed. The samples were polycrystalline with a clear difference in intensity of reflections from crystalline levels for all diffraction peaks. The preferred reflection was (311) plane at $2\theta = 36.92^\circ$, with several secondary reflections of (220), (222), (400), (511) and (440) at $2\theta = 31.27^\circ$, 38.32° , 44° , 60.12° and 56.19° respectively. These results assured predominant of the cubic phase with no reflections for any phases concerning manganese. These peaks were in a perfect match with JCPDS data (JCPDS Card No. 43-1003) [28]. Note that the effect doping concentrations increasing up to 3% did not change the phase of cobalt oxide, and these were the most important conditions of doping, as well as through the alteration in intensity of scattering for the peaks produced. The doping of ions with a smaller ionic radius Mn^{2+} (0.80 Å) than the host material Co^{2+} (0.72 Å) led to a change in the values of lattice constant which could be followed by a deviation in the angle of reflection of the preferred plane towards the smaller angle. A noticeable decrease in the intensity of the preferred reflection (311) plane as shown in Fig. 1(d), (e) and (f) indicated an increase in the lattice constant.

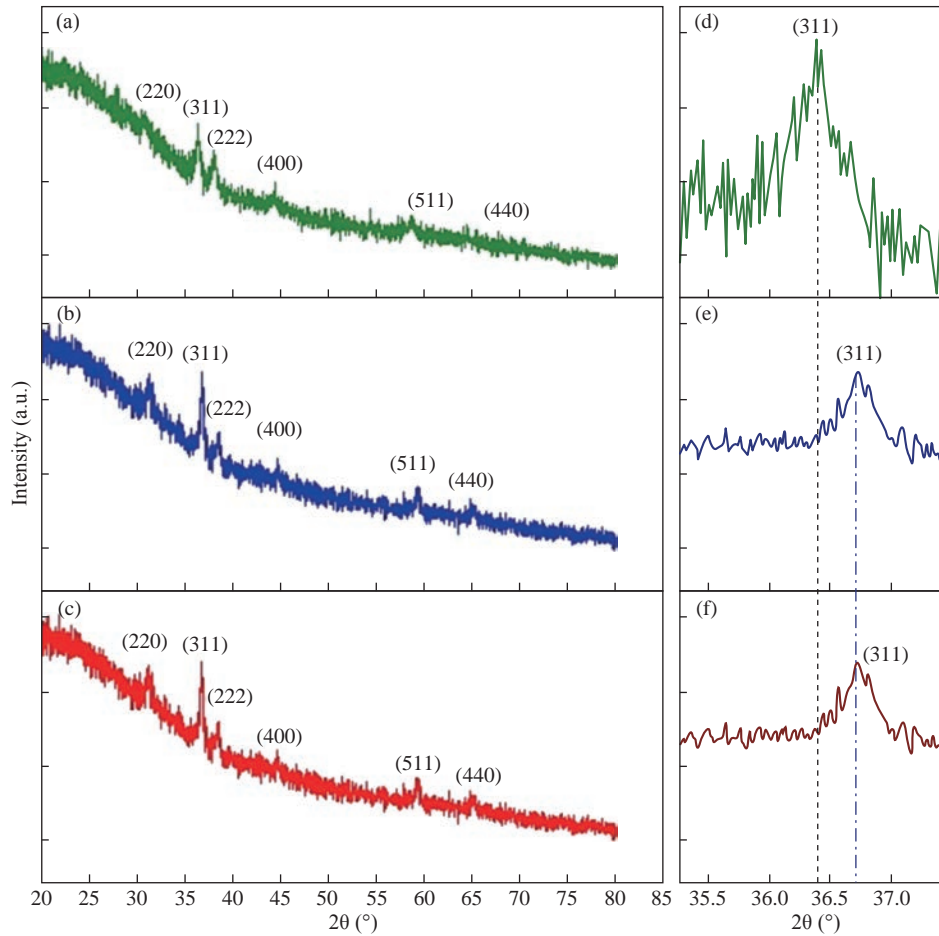


Fig. 1 XRD patterns of (a) pure Co_3O_4 , (b) 1% and (c) 3% Mn doped Co_3O_4 , (d)-(f) drift in 2θ of the preferred plane of pure, 1% and 3% Mn doped Co_3O_4 , respectively.

Fig. 2 offers the crystallite size of the prepared films considered from the FWHM with represented the full width of high maximum; (β) for higher intensity (311) plane by applying Scherrer's Equation (1) [29] showed a decrease in crystallite size (D) via an increase in doping ratio due to interstitial doping processes. The results are shown in Table 1.

$$D = (k\lambda)/(\beta \cos \theta), \tag{1}$$

where λ is x-ray wavelength, θ is Bragg's angle and K is the shape factor.

The Microstrain was calculated from Equation (2) [30, 31], resulting from the growth of the film, which in turn produced pressure and expansion in the

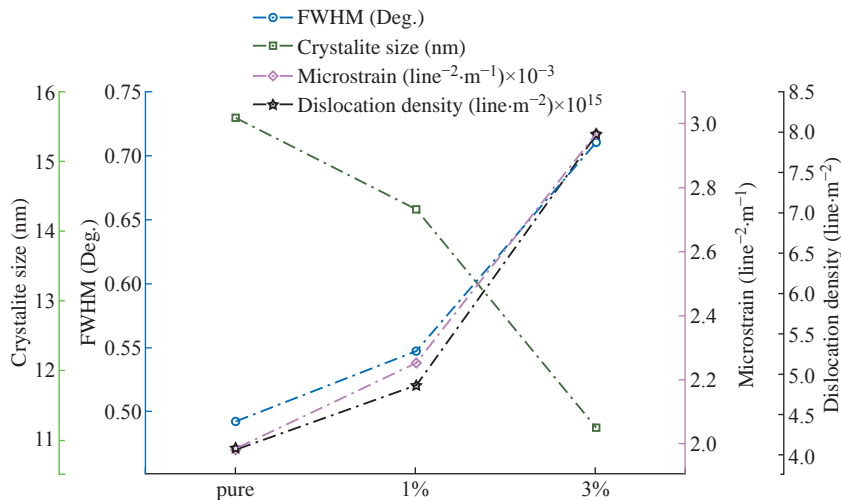


Fig. 2 β , D , ϵ and δ of pure, 1% and 3% Mn doped Co_3O_4 .

lattice. Varying the displacement of the atom based on their original location increased the microstrain ϵ . Fig. 2 shows the increase in doping ratio leading to an increase in microstrain, which was caused by the decrease in crystallinity of the film.

$$\epsilon = \beta/4 \cos \theta, \quad (2)$$

Equation (3) was used to calculate the dislocation density δ ; Fig. 2 displays the dislocation density

increasing as Mn concentration increased, which agreed with the results obtained in previous studies [32, 33].

$$\delta = 1/D^2, \quad (3)$$

The results of ϵ and δ values are listed in Table 1.

Topography of pure and Mn-doped Co_3O_4 films was calculated by AFM (Fig. 3). The roughness, RMS and average grain size were also collected from 2D images.

Table 1 Structural factors of Co_3O_4 : Mn thin films prepared SPT

Sample	Plane (hkl)	2θ ($^\circ$)	Lattice constant A (\AA)	B (Rad)	D (nm)	ϵ ($\text{line}^{-2} \cdot \text{m}^{-1} \cdot 10^{-3}$)	δ ($\text{line} \cdot \text{m}^{-2} \cdot 10^{15}$)
Pure	(311)	36.92	8.98	0.492	15.62	1.98	4.10
1%	(311)	37.02	8.121	0.547	14.32	2.25	4.88
3%	(311)	37.12	8.123	0.711	11.20	2.97	7.97

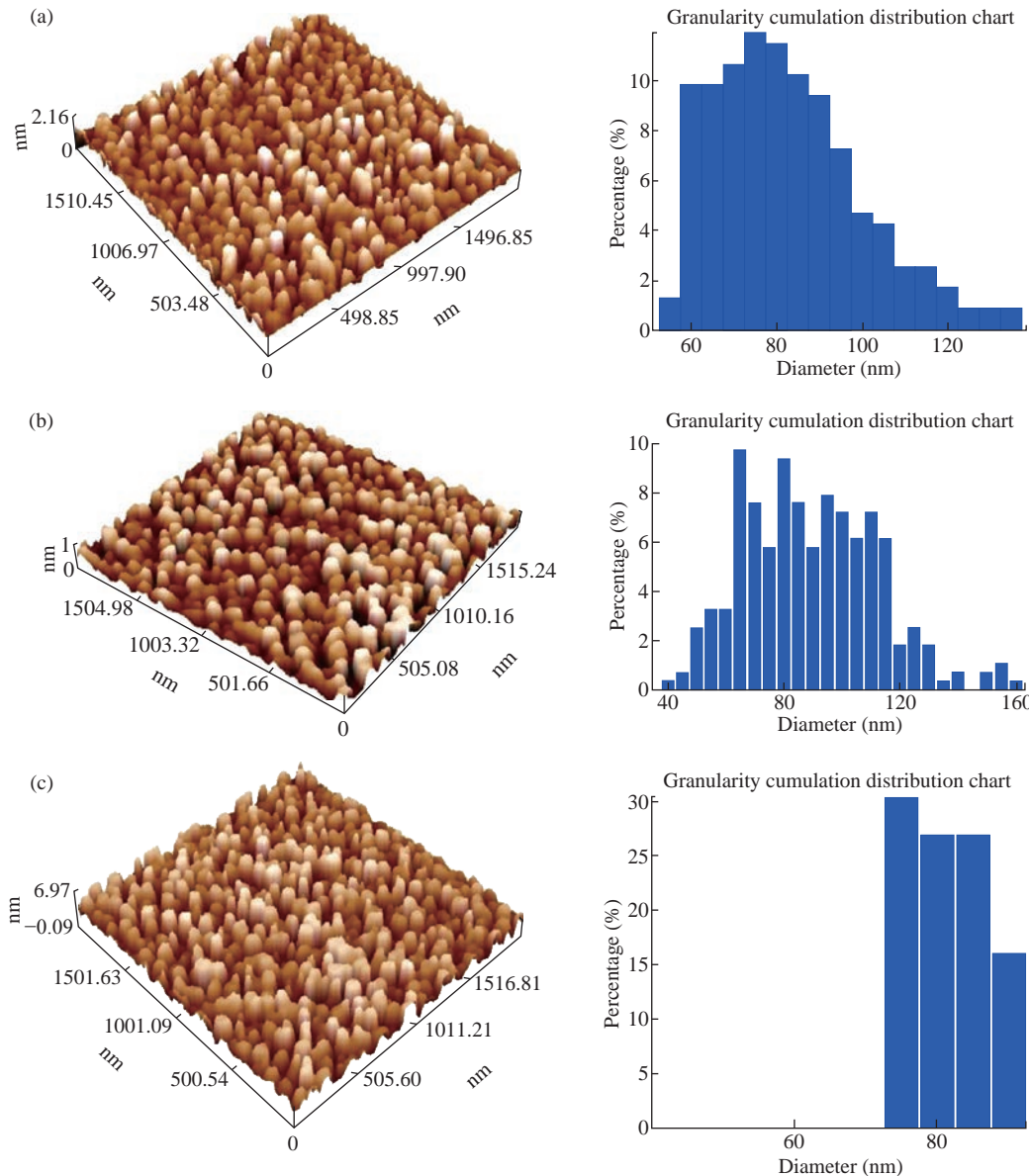


Fig. 3 AFM image of (a) pure Co_3O_4 , (b) 1% and (c) 3% Mn doped Co_3O_4 .

In Fig. 3, a clear variation in morphology of Mn-doped Co_3O_4 films as Mn dopant concentration was increased. The AFM images of the pure Co_3O_4 showed that the particle size was regular in shape with a maximum RMS roughness of 0.431 nm. RMS roughness acts as the standard deviation, which is important to describe surface roughness by statistical methods. As the Mn dopant increased from 1% to 3% (Fig. 3(b) and (c)), the columnar growth enclosed grains with a rod profile could be affected by the merger of Mn on the developing surface. The spreading of the crystallites on surface morphology was uniform with decreases in the RMS from 1.36 nm to 1.23 nm as the Mn concentration increased from 1% to 3%. Apart from that, the average diameter decreased from 86 nm to 76 nm due to decomposition in the lattice.

The optical transmittance of undoped and Mn doped Co_3O_4 films was measured by UV-Vis spectrophotometer. The first beam passed through the undoubted substrate while the second beam passed through the thin film. The transmittance depended strongly on Mn concentration as shown in Fig. 4. The transmittance of the undoped Co_3O_4 film approached 48% in the near-IR spectrum. Fig. 4 demonstrated lower transmittance of 38% to 33% with the increase of doping concentration from 1% to 3%, respectively. The decrease in surface roughness with increasing doping concentration is shown in Fig. 4(a)-(c). Fig. 4(d) represents the absorption values that were observed

to increase with increasing doping concentration that could be related to the doped thin films that could absorb light in a broader range of wavelengths and operate extra light energy than Co_3O_4 thin film. The absorption spectrum took exponential decay with increasing wavelength due to the low energy of the falling photons and their inability to lift electrons from the valence band to the conduction band (Fig. 4(d)).

The values of the energy gap depended on the precise structure of the deposition membrane on the arrangement and distribution of atoms in the crystalline bond [34]. Cobalt oxide compounds have a direct and indirect energy gap [35] and may vary depending on the type of deposition or conditions. Fig. 5(a)-(c) shows the graph of $(\alpha h\nu)^2$ vs E (eV) for direct energy bandgap of pure and Mn-doped Co_3O_4 thin film. The value obtained for the energy gap of the cobalt oxide of 1.435 eV differed from the standard value of 1.419 eV [36]. This difference in the value of energy gap could be achieved by deposition mechanisms or conditions accompanying this deposition. Fig. 5(a) and (b) shows a decrease in the energy gap values from 1.419 eV to 1.367 eV with the increase of Mn concentration from 1% to 3%, respectively. This might be due to a decrease in crystallite size values calculated from the X-ray diffraction pattern and the atomic force microscope, which led to crystallization weakness [37].

Absorption coefficient (α) can be defined through Equation (4). α decreased with increases of λ

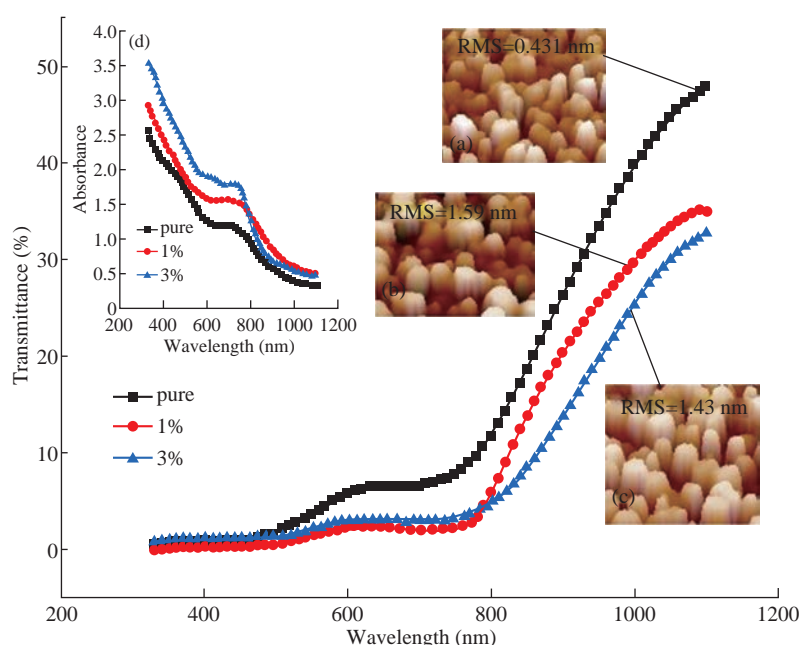


Fig. 4 Optical spectrum transmittance of Co_3O_4 : Mn: (a)-(c) 3D image for pure, 1% and 3% Mn doped Co_3O_4 , respectively; (d) spectrum absorption of Co_3O_4 : Mn.

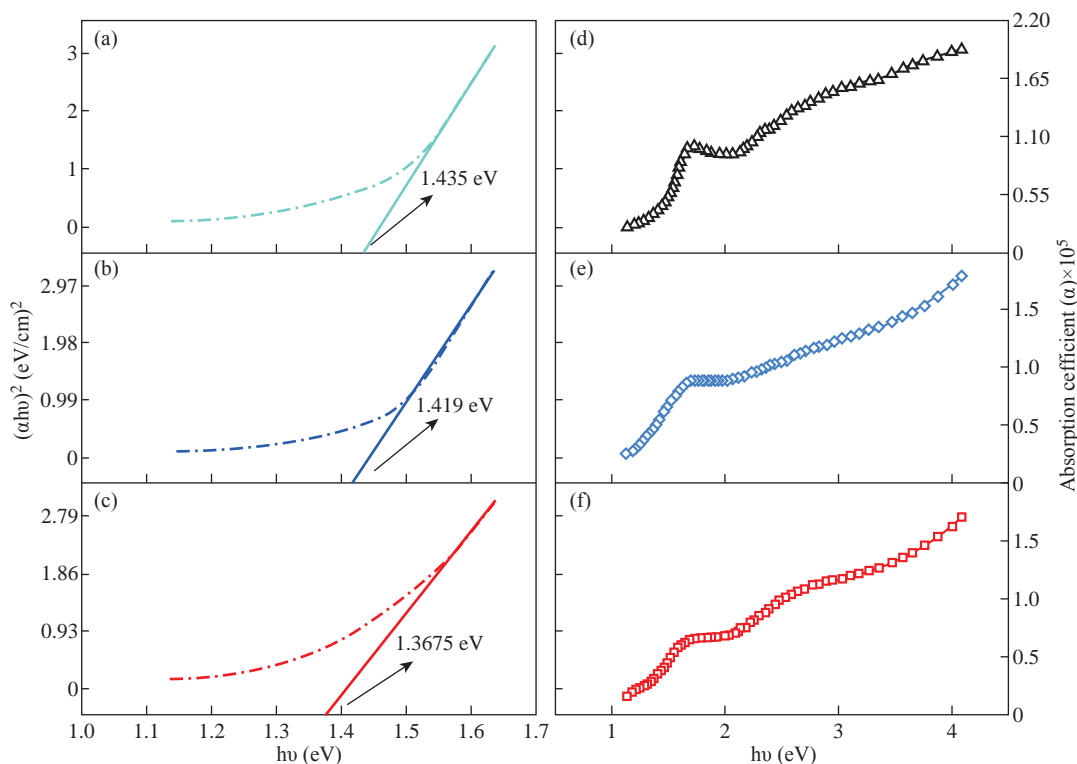


Fig. 5 (a)-(c) The optical energy gap of pure, 1% and 3% Mn doped Co_3O_4 , respectively; (d)-(f) the absorption coefficient of pure, 1% and 3% Mn doped Co_3O_4 , respectively.

increases (Fig. 5(d)-(f)) due to an increase in the Mn concentration leading to an increase in Mn^{2+} ions [38]. α was obtained utilizing the following Equation (4) [39]:

$$\alpha = 2.303A/t \text{ (cm}^{-1}\text{)}, \quad (4)$$

where t is the film thickness.

The refractive index (n) is shown in Fig. 6, calculated from reflectance (R) via Equation (5) [40]. The study of refractive index obtained a clear picture of

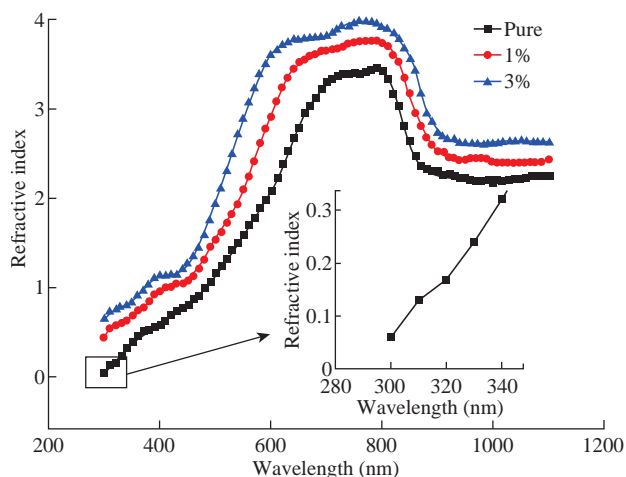


Fig. 6 The refractive index of pure and Mn doped Co_3O_4 , deposited by the spray pyrolysis technique.

the properties and visual behavior of the films recorded for the pure film's refractive index values of 2.6 which were taken down to 2.4 then to 2.3 with an increase in dopant concentration from 1% to 3%, respectively, due to the increased density of the medium, besides the scattering of the incident light at the grain boundaries [41].

$$n = (1 + \sqrt{R}) / (1 - \sqrt{R}). \quad (5)$$

Fig. 7 represents the dynamic resistance change as a function of time for several NO_2 gas concentrations (200, 400, 600, and 800 ppm) at operating temperature of 200 °C for Co_3O_4 and Mn doped Co_3O_4 . It was noted that the resistance values were limited to 5-45 k Ω .m; the greatest resistance was recorded in the presence of air R_a for the undoped samples; the resistance values decreased with the doping ratio due to increasing number of charge carriers. The resistance of the sample decreased with increasing gas concentration from 200 to 800 ppm, which can be due to surface roughness. All recorded samples had a negative connection feature, a decrease in resistance with an increase in concentration of NO_2 gas. When O_2 molecules are adsorbed on the surface of metal oxides (Fig. 7(b)), which leads to the capture of electrons from the conduction band (E_c), the electrons are held on the surface of the films

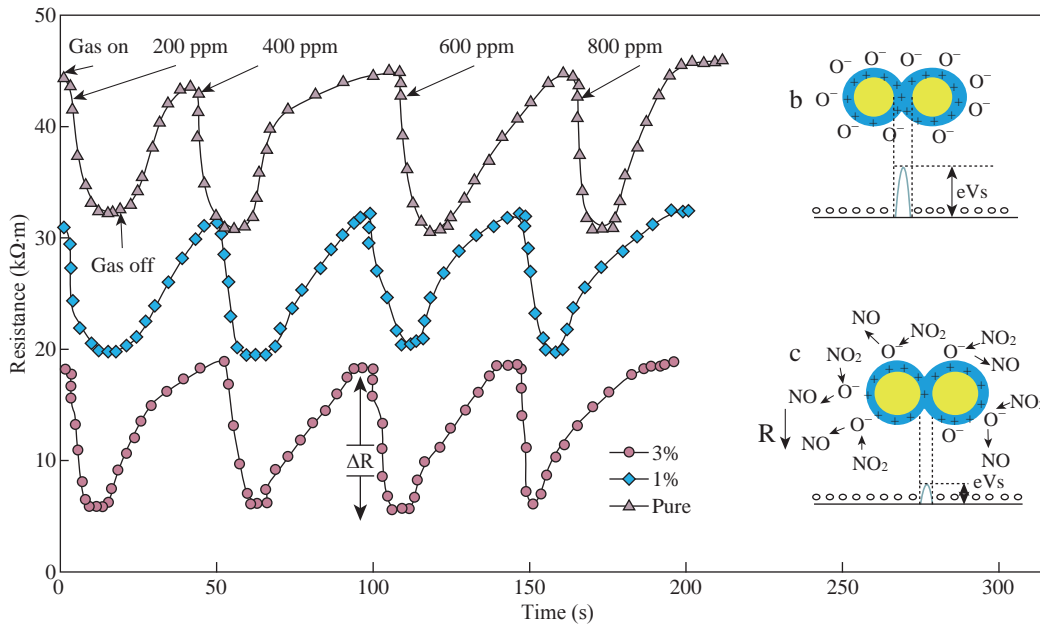


Fig. 7 Resistance change as a function of time of $\text{Co}_3\text{O}_4:\text{Mn}$ films for changed concentrations of NO_2 gas at a working temperature $150\text{ }^\circ\text{C}$.

as models (O^- , O^{2-} , O_2^-). This will raise an electron-depleted region. The electron-depleted region is called the space-charge layer, of which thickness is the length of the band bending region. The interaction of these components with oxidizing gas or competitive adsorption and replacement of the adsorbed oxygen by other molecules decreases, and can reverse the band bending, resulting in a decrease in depletion region of the $\text{Co}_3\text{O}_4:\text{Mn}$ gas sensor as reacting with NO_2 gas and extend in the depletion layer as $\text{Co}_3\text{O}_4:\text{Mn}$ reacting with reduced gas Fig. 7(c).

Fig. 8 represents the sensitivity percentage values

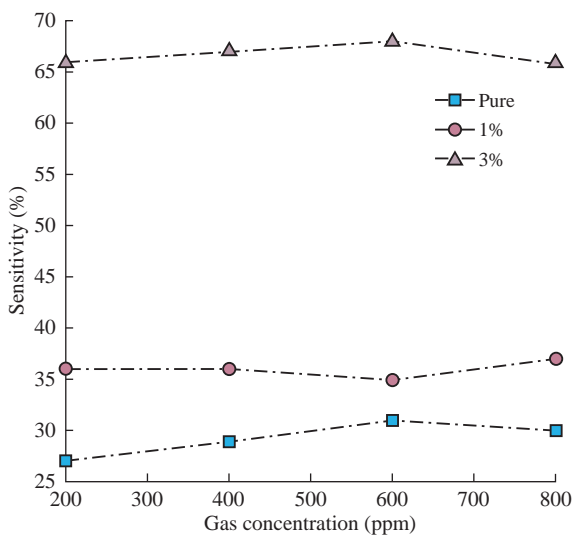


Fig. 8 Percentage sensitivity change of $\text{Co}_3\text{O}_4:\text{Mn}$ films for changed concentrations of NO_2 gas at a working temperature of $150\text{ }^\circ\text{C}$.

calculated from Equation (6) [42]. Fig. 7 depicts $\text{Co}_3\text{O}_4:\text{Mn}$ films that deposited on the glass by the SPT method at an operating temperature of $200\text{ }^\circ\text{C}$ as a function of NO_2 gas concentrations. It could be seen that the highest percentage sensitivity was sample doped with Mn of 3%, which was about 65%, possibly due to the increase in the number of charge carriers of holes owing to the doping and the increase in crystallite size, reducing the depletion layer and potential barrier.

$$\text{Sensitivity} = \Delta R/R_a = |(R_a - R_g)/R_a| \times 100\%, \quad (6)$$

where R_a is resistance of the film sensor in the presence of air, and R_g resistance of the film in NO_2 gas.

Fig. 9(a) represents the response time as a function of gas concentration of NO_2 . It was observed the minimum response time was in the 3% doping concentration, and at the concentration of the gas 800 ppm, it was about 4 s. (Reducing time is a good feature of sensors.) In general, the response time decreased with the increase of gas concentration, which could be due to the increased charge carrier concentration by adsorption and absorption of the gas to the surface.

Fig. 9(b) represents the recovery time or return time as a function of NO_2 gas concentrations, where the minimum recovery time of the 1% model was equal to 10 s (Reducing time is a good feature of sensors). This occurred due to the deforming of films at $150\text{ }^\circ\text{C}$. Continuous exposure to operating temperature caused the film properties to change with the change of gas sensing properties [43].

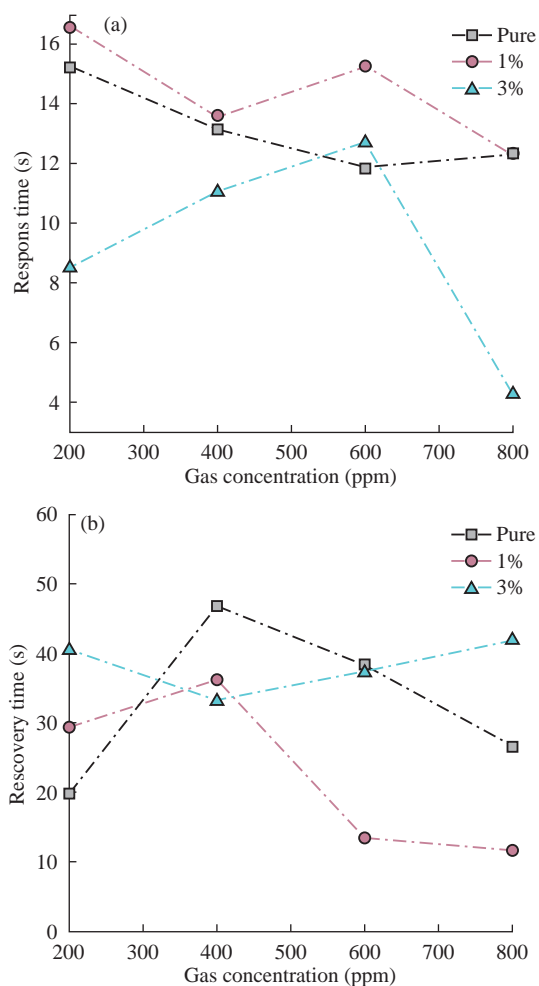


Fig. 9 (a) Response time and (b) recovery time. Change of Co_3O_4 :Mn films for different concentrations for NO_2 gas at a working temperature 150°C .

Conclusions

Undoped and Mn-doped Co_3O_4 were produced by the SPT. The structural description of the samples studied by XRD displayed polycrystalline with cubic structure; the crystallite size reduced from 15.62 nm for the pure Co_3O_4 to 11.2 nm upon 3% Mn-doped Co_3O_4 samples. The morphology investigation by AFM showed a columnar growth containing grains with a rod shape, which can be caused by a combination of Mn on the developing surface. The incorporation of Mn into Co_3O_4 weakened the energy band due to the formation of surface-associated defects in the nanoparticles. We concluded that the electrical resistance of the films were reduced from 45 k Ω to 18 k Ω with the increasing of doping rate, and that the best sensitivity to the gas NO_2 was 65% by 3% Mn content. With the concentration of gas 800 ppm at a temperature of 150°C , the best recovery time was 10 s, and the best response time was 4 s to the doped film by 3%,

with the concentration of gas 800 ppm at a temperature of 150°C .

Acknowledgments

This study was completed with the help of Mustansiriyah University, College of Science, physics department, and college of Education, physics department.

Conflict of Interests

The authors declare that no competing interest exists.

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