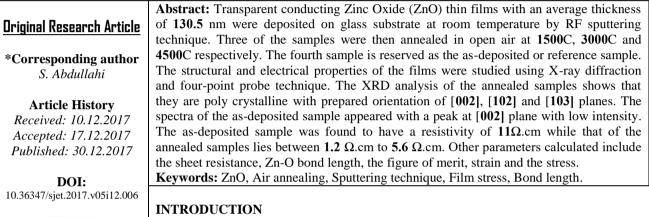
Scholars Journal of Engineering and Technology (SJET) *Abbreviated Key Title: Sch. J. Eng. Tech.* ©Scholars Academic and Scientific Publisher A Unit of Scholars Academic and Scientific Society, India www.saspublishers.com

The Role of Air Annealing On the Structural and Electrical Properties of Zinc Oxide (Zno) Thin Film Deposited By Rf Sputtering Technique

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Presently, nano crystalline materials in the form of thin films have opened new chapter in the field of electronic applications, since material properties could be changed by changing the crystallite size and/or thickness of the film. Development of such materials, whose structural and electrical properties could be controlled, will be useful in many ways. For example, optoelectronic devices, particularly solar energy conversion devices could be modified accordingly. The synthesis of binary metal chalcogenide of groups II–VI semiconductors such as ZnO in a nano crystalline form has been a rapidly growing area of research due to their important non-linear optical properties, luminescent properties, quantum-size effect and other important physical and chemical properties.

The properties of materials prepared by different methods are critically dependent on the nature of preparation technique and subsequent heat treatments like annealing in air, vacuum or different gaseous environments like H₂, N₂, Ar, etc.

Zinc oxide (ZnO) is an inexpensive strategic material for various photonic applications. As one of the most important binary II-VI compounds, ZnO is a direct semiconductor of wurtzite structures that is short wavelength suitable for optoelectronic applications [1, 2]. ZnO exhibits a direct bandgap of 3.3 to 3.37 eV at room temperature and efficient radiative recombination. Its large exciton binding energy of 60 meV which is 2.4 times higher than that of gallium nitride (GaN), makes it transparent in visible light and operates in the UV to blue wavelengths, paves the way for an intense near band edge excitonic emission at

room and even higher temperatures [3, 4]. ZnO thin films presents many remarkable characteristics due to their large bond strength, good optical quality, extreme stability of excitons and excellent piezoelectric properties. For this reason, ZnO thin films have many potential applications in various technological domains such as transparent conducting films/electrodes in display devices and solar energy cells, surface and bulk acoustic wave devices (SAW) and acoustic-optical devices, light emitting diodes (LEDs) and laser diodes (LDs).

ZnO thin films have been prepared by a variety of deposition techniques such as rf sputtering [5-8], Spin coating [9], Spray pyrolysis [10], Electrochemical deposition [11, 12], Sol-gel [13, 14], Thermal evaporation [15], SILAR [16, 17] and Aqueous Chemical Growth method [18].

Available online at <u>https://saspublishers.com/journal/sjet/home</u> 704 In this contribution, thin films of ZnO were deposited on glass substrate by rf sputtering technique. Three of the four samples were annealed in open air to analyze their structural electrical properties. Sputtering offers a wide range of advantages, such as easy adaptation to large-scale and reproducible manufacturing. After all annealing is known to improve crystalline qualities conductivity of thin films [4].

MATERIALS AND METHODS

Substrate cleaning

Commercially available corning glass 7059 micro slides of dimensions 25.4mm ×76.2mm × 1.2 mm were boiled in chromic acid for 2 hours at 100°C in an ultrasonic bath, boiled in dilute hydrogen peroxide solution for 15 minutes, rinsed in acetone, cleaned with double distilled water and finally dried with flowing Nitrogen gas and placed in the sputtering chamber.

ZnO thin film formation

After evacuating the chamber to a base vacuum of 1×10^{-5} torr, argon/oxygen of (4N purity) gas mixture was fed to the system. The ZnO target (99.99% pure) of 4 cm in diameter was sputtered in pure argon/oxygen atmosphere for about 5 minutes to remove the surface layer of the target before the deposition of the film. ZnO films were deposited on the glass substrates at room temperature to a film thickness of 130.5 nm. The RF power used was 60W, target to substrate distance was constantly kept at 7cm. This deposition was carried out under argon/oxygen gas flow of 10 sccm.

There were four samples from which 3 were annealed in open air for 60 minutes at 150° C, 300° C and 450° C. The fourth sample was reserved as the reference or as-deposited sample.

Characterization of ZnO thin films

The thickness of the deposited films was measured using Veeco Dektak 150 profilemeter. The crystal structure of the films was inspected using an X-ray diffraction performed in $2\theta/\omega$ at a voltage of 45kv and a current of 40 mA. The sweeping angle is 20 to 80 degrees, the scan speed is 0.8 degrees/minute at a scan step of 0.02, employing a *Cu ka tube* ($\lambda = 0.1540598 nm$) radiation. The electrical analysis of the samples was carried out using four-point probe.

RESULTS AND DISCUSSION XRD studies

Figure 1 shows XRD pattern for the asdeposited as well as the annealed samples. This XRD spectra reveal that all the annealed thin films were polycrystalline with hexagonal structure (JCPDS 36-1451) [19]. There are no visible peaks for the asdeposited sample except that of peak [002] which appeared with low intensity. An improvement in the XRD structure can be noticed in the structure of the annealed samples. This indicates that annealing can enhance the structure of thin film as sizes of grains on the surface increases [20]. The diffraction peaks of the annealed samples are indexed as (002) plane at 2θ = 34.02° , 34.38° and 34.43° respectively. The peak position and the intensity are in accordance with the JCPDS card number #01-074- 9942. The XRD patterns of all the annealed samples indicated enhanced preferred orientation along the c-axis.

The grain size was calculated using Debye-Scherrer's formula [21, 22] given by Eq. (1).

$$d = \frac{0.9\lambda}{\beta\cos\theta} \tag{1}$$

where "d" is the grain size or crystallite size, ' λ ' the wavelength of the X-rays (1.5405Å), ' θ ' the diffraction angle, and β is the full width at half maximum of (002) peak of the XRD data.

The lattice constants a and c, for a hexagonal structure are calculated by [16] using Eq. (2);

$$d_{hkl}^2 = \left(\frac{4(h^2 + k^2 + hk)^2}{3a^2} + \frac{l^2}{c^2}\right)^{-1}$$
(2)

where d is the interplanar space and h, k, l are Miller indices.

The strain (ϵ) and stress (σ) in the film, along the c-axis were found to be within an acceptable range by using Eq. (3) and Eq. (4) [22].

$$\varepsilon = \frac{c_{film} - c_{bulk}}{c_{bulk}} \tag{3}$$

$$\sigma = 2.33 \times 10^{11} \left(\frac{\sigma_{flim} - \sigma_{bulk}}{\sigma_{bulk}}\right) \tag{4}$$

where c_{film} and c_{bulk} (5.2 Å) are the lattice parameters of the film and unstrained ZnO respectively.

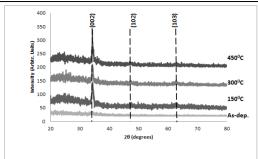


Fig-1: XRD pattern for as-deposited and the annealed samples

The calculated lattice constants are given in table 1. These lattice constants are seen to be in an acceptable range compared with those measured by [12-14] and also in agreement with lattice constants of ZnO Zincite phase existing in the literature of ASTM card: a =3.2648Å and c =5.2194Å. The calculated d-spacing is also within the range of JCPDS data (36-1451) for (002) planes given as 2.6049.

The slight variations seen in d-spacing could be due to the interstitial zinc and it suggests that the unit cell might be elongated along the c-axis, and there is stress in the plane. The FWHM of the as-deposited sample stood at 0.5760. The FWHM increases with increase in the annealing temperature for the annealed samples. The grain size (D) for the as-deposited sample is 2.5199. Grain size increases with increase in the annealing temperature for the annealed samples. An increase in the FWHM indicates a degradation in the crystal quality [23]. The Zn-O bond length was also calculated [22] using Eq. (5) and Eq. (6).

$$L = \sqrt{\left(\frac{a^2}{3} + \left(\frac{1}{2} - u\right)^2\right)c^2}$$
(5)
$$u = \frac{a^2}{3c^2} + \frac{1}{4}$$
(6)

Sample	FWHM(20)	Crystallite	2 0 (002)	d-	Lattice	Bond	$Strain(\epsilon)$	Stress(σ)
		size (nm)	Peak	spacing	parameters a	Length	$\times 10^{-2}$	Gpa
			only		and c (Å)	(L)(Å)		
As-	0.5760	129.82	34.40°	2.60	a=3.18 c=5.18	2.11	-0.38	8.9
deposited								
Annealed	1.1520	61.94	34.02°	2.63	a=3.23 c=5.10	2.15	-0.19	4.5
at 150 ⁰ C								
Annealed at 300 ⁰ C	0.6720	109.77	34.38°	2.61	a=3.19 c=5.15	2.13	-0.96	2.2
Annealed	0.4800	158.92	34.43°	2.60	a=3.18 c=5.20	2.14	0	0
at 450°C	011000	100072	00	2.00	a 2.110 C 2.20		Ũ	Ũ

Table-1: Grain size and FWHM for as-deposited sample and samples annealed in air

Electrical resistivity and sheet resistance

Table 2 shows the resistivity and the sheet resistance of the ZnO thin films deposited by RF sputtering technique. According to the table, the asdeposited sample exhibits the highest resistivity of 11 Ω .cm. The resistivity of the annealed samples varies with the annealing temperature. Decrease in resistivity for the annealed samples as compared with the asdeposited samples indicates the semiconducting nature of the samples. Resistivity of the TCO film depends strongly on the chamber oxygen pressure as well as the stoichiometry of the film. The electrical resistivity of thin films may also be affected by isotropic background scattering (arising from phonons and point defects), external surface scattering and grain boundary scattering [22].

The sheet resistance of the samples is also

Available online at <u>https://saspublishers.com/journal/sjet/home</u> 706 displayed in table 2. The as-deposited sample shows the highest sheet resistance of 4.99 Ω . Sheet resistance varies for the annealed samples. The high sheet resistance observed for the as-deposited sample and the sample annealed at 150°C could be attributed to surface scattering and the decrease in carrier concentration [24]. For these films, more defects act as scattering centers which results in the formation of trapping states capable of trapping carriers and thereby immobilizing them. This reduces the number of free carriers available for electrical conduction.

It has also been suggested that a decrease in sheet resistance with respect to increase in annealing temperature may be due to the better crystal orientation of the sample [25]. It is known that undoped ZnO thin films generally exhibit n- type conductivity, the value of which depends on the deposition parameters [26]. The suitability of transparent conducting oxides (TCOs) for optoelectronic applications can be quantified by a factor known as the figure of merit (ϕ) which is a combination of high electrical conductivity and low absorption (reported elsewhere) of visible light, calculated using Eq. (7) [23]. The as-deposited sample

exhibited the lowest figure of merit while the sample annealed at 300° C exhibited the highest.

 $\oint \frac{1}{\alpha \rho} \tag{7}$

where α is the absorption coefficient at 550 nm and ρ the electrical resistivity.

Table-2: Sheet resistance (R_s) and resistivity of as-deposited sample and samples annealed in air

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Annealing Temperature(⁰ C)	Resistivity (Ω.cm)	Sheet resistance $(R_s)\Omega$	Figure of merit $(\phi)(\Omega^{-1})$					
As-deposited	11.0	4.99	0.99×10^{-6}					
150	5.6	2.27	2.05×10^{-6}					
300	1.2	0.544	3.29×10^{-6}					
450	1.8	0.816	1.5×10^{-6}					

CONCLUSIONS

From the above studies, it is concluded that the ZnO thin films deposited by RF sputtering technique at room temperature and annealed at various temperatures have wurtzite crystalline structure with preferential orientation along the c-axis. The magnitude of the figure of merit shows that these films can be used for optoelectronic applications.

ACKNOWLEDGEMENTS

The authors are indebted to the World Bank through the Science and Technology Post Basic (STEP-B) of the Usmanu Danfodiyo University Sokoto.

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