Structural, Optical Properties

by Johansah Liman

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Structural, Optical Properties and Raman Spectroscopy of In₂O₃ Doped LiTaO₃ Thin Films

Nani Djohan^{1*}, Budi Harsono¹, Johansah Liman¹, Hendradi Hardhienata² and Irzaman^{2*}

- Department of Electrical Engineering, Faculty of Engineering and Computer Science, Universitas Kristen Krida Wacana, Jl. Tj. Duren Raya No. 4, Jakarta 11470, Indonesia
- ² Department of Physics, Faculty of Mathematics and Natural Sciences, Bogor Agricultural University, Jl. Raya Dramaga, Bogor 16680, Indonesia

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ABSTRACT

In this experiment, undoped, 2 wt.%, 4 wt.% and 6 wt.% In_2O_3 doped LiTaO₃ thin films were successfully prepared by utilizing a spin 11 ter to carry out chemical solution deposition on the substrate surface (CSD method). The films were grown on the p-type Si (100) substrates with 2 M in 2-methoxyethanol precursor, whose solubility was twisted at 4000 rpm for 30 seconds. Crystalline formation of the films was carried out at annealing temperature 850 °C, held for 15 hours at a temperature rise rate of 1.67 °C/min. In term of XRD analysis, the structural properties of LiTaO₃ thin film undergo increment in crystallite size and lattice 13 ameter values as the concentration of indium doping increase. The optical properties and Raman spectra of the films were then obtained using UV-Vis spectrometer and Raman spectroscopy. From the XRD measurement, the result shows a hexagonal crystal structure with lattice parameters a = 5.032-5.051 Å and c = 13.643-13.676 Å, and from the UV-Vis data, we observed that the films have a = 5.034-5.184 Ev energy gap with a = 1.70364373 - 1.70364377 refractive index. Raman analysis produces peaks of LiTaO₃, a = 1.70061273 and a = 1.70061273 a

Keywords: LiTaO₃, In₂O₃, x-ray diffraction, Raman spectroscopy, uv-vis spectroscopy

1. INTRODUCTION

A thin film made from Lithium tantalate (LiTaO₃) is an essential optical material, mainly due to its excellent electro/acousto-optical properties, high electro-optic and pyroelectric coefficient of 30.5 pm/V and 230 μ C,m⁻².s⁻¹ respectively, and high curie temperature of 655 °C [1,2]. The thickness of the sensitive cells is inversely proportional to the detector response [36]. LiTaO₃ is very attractive and promising material for optical sensor [3,7]. The performance of thin films can be increased by the addition of doping materials. The concentration of the doping materials affects the structural and optical properties of the thin films. Mendoza 18 deposited Fedoped LiTaO₃ thin films by magnetron sputtering and they investigated the effect of iron doping on the crystalline structure. Irzam 12 et al. [14] prepared lanthanum doped LiTaO₃ thin films by CSD method and they investigated the effect of dopant addition on the crystal formation and the optical properties of the films.

There are various kinds of technology to synthesize $LiTaO_3$ thin films. X-ray Diffraction (XRD) technology is based on diffraction of X-Ray when scattering light with a wavelength passing through a crystal lattice with a distance between crystal fields. Previous study stated on the crystal structure and lattice parameter found that the diffraction angle depends on the gap of the

^{*}Corresponding author: nani.djohan@ukrida.ac. irzaman@apps.ipb.ac.id

lattice width, thus affecting the diffraction pattern [12]. In contrast, the diffraction light intensity depends on the number of crystal lattices having the same orientation. Crystal system, parameter of the lattice, the degree of crystallinity, structure type, and orientation can be determined by this method. LiTaO $_3$ thin film with high purity and best intensity was selected from the XRD measurement [3,9,10].

Solid-state and molecule vibrational properties can be analyzed by using Raman spectroscopy [11,12]; thus, strain [12,15], doping [12,16,17], stoichiometry [12,13,14] and crystallinity property information can be obtained by this technology. Density functional theory (DFT) is the combination of Raman with theoretical approaches yielding phonon eigenvectors [12]. In this experiment, four types of solution (LiTaO3, LiTaO3 + 2 wt.% In_2O_3 , LiTaO3 + 4 wt.% In_2O_3 , and LiTaO3 + 6 wt.% In_2O_3) were produced and structural, optical properties and raman spectra were carried out.

2. MATERIAL AND METHODS

The p-type Si (100) substrates cut in size of 1x1 cm² were ultrasonically cleaned in acetone (C_3H_6O) and repeated sequentially using methanol (CH_3OH) and deionized water successively for 15 minutes [5,9,24-27]. The undoped LiTaO₃ solution was prepared by dissolving 0.5897 gram of lithium tantalate (\geq 99.99%, Sigma-Aldrich 704393-5G) in 2.5 ml of 2-methoxyethanol. The 2 wt.%, 4 wt.%, and 6 wt.% In_2O_3 doped LiTaO₃ solutions were prepared by dissolving 0.5897 gram of lithium tantalate and 0.01179 gram, 0.0236 gram, and 0.03538 gram of indium (III) oxide (99.99%, Sigma-Aldrich 289418-10G), respectively in 2.5 ml of 2-methoxyetanol.

To get the homogeneous LiTaO₃ solution, it was stirred using Vortex 3000 mixer and then inicated using an ultrasonicator device (J.P Selecta) for 30 minutes [5,9]. Furthermore, the thin films were deposited on the p-type Si (100) substrates using spin coater for 30 seconds using a rotating speed of 4000 rpm. This process was repeated three times with a one-minute time interval [5,6,9,19,24-29]. The annealing process was done using Nabertherm B410 furnac 4 at temperature 850°C for 15 hours, with a temperature increment speed of 1.67°C/min. The structural properties of the thin-film crystal were then obtained with an X-Ray Diffractometer (Bruker D2 PHASER) with the interval of 2 theta angles from 20 to 80 degrees with a 0.02 degrees step, where thin films have formed crystals at specific angular pezes with hexagonal-shaped crystal structures. Furthermore, the thin films were characterized by using a UV-Vis spectrometer (Ocean Optics USB4000-UV-Vis) and Raman spectroscopy (Horriba iHR550) to investigate the transversal and longitudinal phonon modes of the films [11,12,18,20-23].

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3. RESULTS AND DISCUSSION

3.1 Structural properties

The XRD patterns of LiTaO₃ thin films we determined from $20^{\circ} \le 20 \le 80^{\circ}$ intervals with a scanning rate of 0.02° /minute [9,14,30-32]. Figure 1 shows the XR patterns of undoped and In₂O₃ doped LiTaO₃ thin films [9,14,30]. The Miller index values were assigned by referring the standard x-ray diffraction powder patterns file (JCPDS) Natl Bur, 1977) and were in good agreement with the values reported in the file. From the Rietveld refinement result, the Goodness of Fit (GOF) for the undoped and In₂O₃ doped LiTaO₃ thin films varies between [43]9 and 2.91. The Weighted R profile (R_{wp}), expected R profile (R_{exp}) and GOF are given in Table 1. The XRD patterns showed that LiTaO₃ thin films were both amorphous and crystalline with hexagonal crystal structure. The percentage of amorphous and crystalline are given by the crystallinity index (CI) in the Table 1. All the XRD patterns showed a preferred strong peak at (012) diffraction plane. The peaks profiles of the two most dominant peaks at (012) and (104) diffraction plane was

obtained by Gaussian fitting. Peak position (2θ) and the full width at half maximum (FWHM) from the peaks are shown in Table 2.

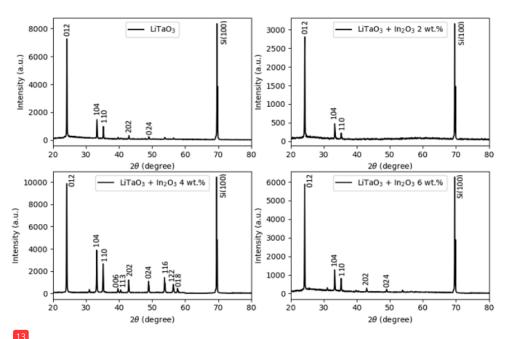


Figure 1. XRD patterns of LiTaO₃ thin films with different doped In₂O₃ at annealing temperature 850°C

The crystallite size (D) of the undoped and In_2O_3 doped $LiTaO_3$ thin films was calculated using Debye Scherer's formula:

$$D = \frac{k\lambda}{\beta cos\theta} \tag{1}$$

where k is the shape factor (= 0.9), λ is the X-ray radiation wavelength (= 0.15406 nm), β is FWHM of diffraction peak in radian, and θ is the Bragg's diffraction angle in radian. The highest crystallite size values observed were 51.408 nm and 46.620 nm for (012) and (104) diffraction plane in 6 wt.% In₂O₃ 30 ped LiTaO₃ thin film, respectively. The crystallite size decreased in line with the decrease in doping concentration, as shown in Table 2. The decrease of crystallite size may be due to better integration of In₂O₃ ion into the LiTaO₃ structure [38].

The dislocation density (δ) of the undoped and In_2O_3 doped LiTaO $_3$ thin films could be obtained using the Williamson-Smallman formula:

$$\delta = \frac{1}{n^2} \tag{2}$$

where D is the crystallite size of the films obtained from Eq. 1. It was observed that the dislocation density decreases as doping concentration increases.

The lattice strain (ϵ) of the undoped and In_2O_3 doped LiTaO₃ thin films was calculated by using the following formula:

$$\varepsilon = \frac{\beta}{4\tan\theta} \tag{3}$$

The data show that when doping concentration increases beyond 2 wt.%, lattice strain will decrease. Throsuggests the creation of fewer defects in the LiTaO₃ lattice. The same finding was obtained by Sahoo et al. [39] in indium doped ZnO thin films. All the calculated results from the XRD data are shown in Table 2.

The interplanar spacing (d_{hkl}) for each peak can be calculated using Bragg's formula, as shown in Eq. 4. The lattice parameters (a and c) can be calculated from the interplanar spacing equation for hexagonal lattice as shown in Eq. 5, where hkl is the Miller indices of the plane of diffraction.

$$d_{hkl} = \frac{\lambda}{2\sin\theta} \tag{4}$$

$$\frac{1}{d_{hkl^2}} = \frac{4}{3} \left(\frac{h^2 + hk + k^2}{a^2} \right) + \frac{l^2}{c^2} \tag{5}$$

The calculated lattice parameters for undoped and various wt.% In_2O_3 doped LiTaO₃ thin films are tabulated in Table 3. From the data in Table 3, the calculated lattice parameters were slightly different from the JCPDS data due to differences in annealing temperature. The data also showed that the lattice parameters increased with increasing doping concentration.

Table 1 R-profile, Goodness of Fit and crystallinity index of LiTaO3 thin films doped with In2O3

Sample	Rwp	Rexp	GOF	Crystallinity Index (%)
LiTaO ₃	25.7	21.8	1.39	68.53
$LiTaO_3 + 2$ wt.% In_2O_3	43.1	28.82	2.24	51.06
$LiTaO_3 + 4$ wt.% In_2O_3	18.7	10.94	2.91	88.98
LiTaO3 + 6 wt.% In2O3	34.4	23.68	2.11	65.32

Table 2 Crystallite size, dislocation density, and lattice strain of LiTaO3 thin films doped with In2O3

Sample	Diffraction plane	2θ (deg)	FWHM (deg)	Crystallite size (nm)	Dislocation density (cm ⁻²) x 10 ¹⁰	Lattice strain x 10 ⁻³
LiTaO ₃	0 1 2	24.186	0.160	50.805	3.874	3.257
	104	33.293	0.197	42.001	5.669	2.881
LiTaO ₃ + 2 wt.% In ₂ O ₃	0 1 2	24.217	0.167	48.638	4.227	3.398
	104	33.333	0.197	42.016	5.665	2.877
LiTaO ₃ + 4 wt.% In ₂ O ₃	012	24.131	0.160	50.685	3.893	3.272
	104	33.231	0.189	43.834	5.204	2.765
LiTaO ₃ + 6 wt.% In ₂ O ₃	0 1 2	24.187	0.158	51.408	3.784	3.218
	104	33.293	0.178	46.620	4.601	2.596

 $\textbf{Table 3} \ \text{Lattice parameter values of LiTaO}_3 \ \text{thin films doped with } In_2O_3 \ \text{hexagonal structure}$

Lattice parameter	LiTaO ₃	LiTaO ₃ + 2 wt.% In ₂ O ₃	LiTaO ₃ + 4 wt.% In ₂ O ₃	LiTaO ₃ + 6 wt.% In ₂ O ₃	JCPDS (Natl Bur, 1977)
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c (Å)	13.658	13.643	13.676	13.658	13.755
a (Å)	5.038	5.032	5.051	5.038	5.153

3.2 Optical properties

The optical properties of the undoped and In_2O_3 doped $LiTaO_3$ thin films were investigated by analyzing the reflectance spectra in the wavelen 23) between 230 nm and 850 nm (UV-Vis range). The reflectance spectra of the synthes 23 d films are shown in Figure 2. In the reflectance spectra, it is observed that 25 e films have low reflectance in the ultra-violet range and high reflectance in the visible range. The band gap energy (Eg) 15 ues of the synthesized films could be estimated from the reflectance spectra. The recorded reflectance spectra were first transformed to the absorption spectra by applying the Kubelka-Munk function (F(R)). Then, the Tauc plot method is employed to determine the band gap energy [5,9,35]. The Kubelka-Munk function and the Tauc formula are given in Eq. 6 and Eq. 7, respectively.

$$F(R) = \frac{\kappa}{S} = \frac{(1-R)^2}{2R} \propto \alpha_{K-M} \tag{6}$$

$$(\alpha_{K-M}hv)^{1/n} = B(hv - Eg) \tag{7}$$

and S are absorption and scattering coefficients, respectively. R is reflectance value, hv is the photon's energy, B is a constant, Eg is the band gap energy, and n is the nature of the electron transition (= ½ for direct transition band gap).

From the Tauc plot, it is found that the optical band gap energy of undoped, 2 wt.%, 4 wt.%, and 6 wt.% In_2O_3 doped $LiTaO_3$ thin films are 5,034 eV, 5,079 eV, 5.112 eV, and 5,184 eV, respectively, as shown in Figure 3. From the data, it was observed that the increase in doping concentration also increase the band gap energy of the films. This result is parallel agreement to Singh et al. [40].

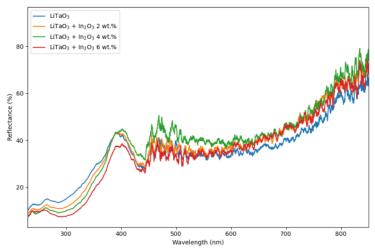


Figure 2. Reflectance spectra of LiTaO₃ thin films

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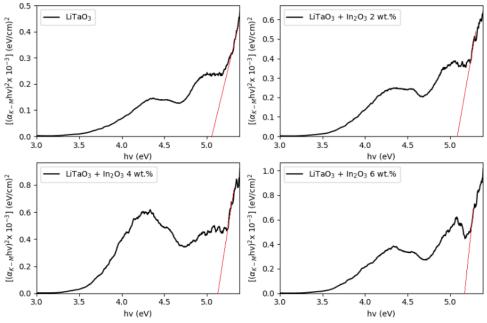


Figure 3. Energy band gap of LiTaO3 thin films

Refractive index (n_e) of the films were obtained by using Sellmeier formula [9,36,37]:

$$n_e^2(\lambda, T) = A + \frac{B + b(T)}{\lambda^2 - [C + c(T)]^2} + \frac{E}{\lambda^2 - F^2} + \frac{G}{\lambda^2 - H^2} + D\lambda^2$$
(8)

where λ is the temperature dependent, and A, B, C, D, E, F, G, H are the band gap energy value, b(T), c(T) is the temperature dependent, and A, B, C, D, E, F, G, H are the band gap energy increased. The data show that the value of the refractive index also increases as the band gap energy increased. The refractive index, band gap energy and related color spectra of undoped, 2 wt.%, 4 wt.%, and 6 wt.% In_2O_3 doped $LiTaO_3$ thin films were listed in Table 4. The result was acceptable as previous study by Zielinska et al. [30].

Table 4	<mark>13</mark> e energy	gap and th	e refractive	index of	LiTaO3	thin film	S

Sample	Band gap (eV)	Wavelength (nm)	Color spectra	Refractive index
LiTa03	5.034	246.855	ultraviolet	1.70364373
LiTa03 + 2 wt.% In203	5.079	244.692	ultraviolet	1.70364374
LiTa03 + 4 wt.% In203	5.122	242.529	ultraviolet	1.70364375
LiTa03 + 6 wt.% In203	5.184	239.715	ultraviolet	1.70364377

3.3 Raman Spectrum

Raman spectrum of LiTaO₃ single crystals with various stoichiometric was used to investigate the compositional uniformity of the crystals [11,12,18]. The first principles, the peaks of Raman intensity were assigned by referring to the mode energy [12,33,34]. The phonons identification may become complicated because some Raman intensity might be lost on the calculated modes [12]. Longitudinal phonon mode was observed when polarization vectors and phonon wave

vector are parallel, whereas transversal phonon mode was observed when polarization vectors and phonon wave vector are perpendiculars.

Figure 4 shows the Raman spectrum 17 indoped, 2 wt.%, 4 wt.%, and 6wt.% In_2O_3 doped $LiTaO_3$ thin films. In this experiment, Raman spectra were obtained by Raman spectroscopy in the range of 100 to 1000 cm⁻¹.

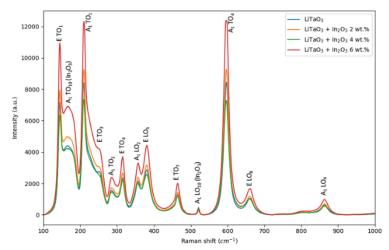


Figure 4. Raman spectroscopy of LiTaO3

Table 5 shows the result of active phonon frequencies of undoped, 2%, 4%, and 6% In ${}_2O_3$ doped LiTaO $_3$ thin films. The experiment results show that the most significant deviation is 5 cm $^{-1}$ at A $_1$ TO $_4$ and the mean deviation is 1.82 cm $^{-1}$.

Table 5 Theory and experiment result of active phonon frequencies of LiTaO $_3$

Symmetry	Mode	Exp.	Literature [9]	Mode	Exp.	Literature [9]
A1	TO1	209	209	LO1		255
A1	TO2	285	286	212	356	355
A1	тоз		376	LO3		403
A1	TO4	596	591	LO4	863	866
E	TO1	144	144	LO1		190
E	TO2		199	LO ₂		
E	TO3	253	253	LO3		279
E	TO4	315	319	LO4		
E	TO5		409	LO5	380	381
E	T06		420	LO6		453
E	TO7	463	459	LO7		
E	T08		590	LO8	661	660
E	TO9		669	LO9		866
A1	TO10	165		LO10	521	

Phonon activities in hexagonal crystal structures (XRD analysis) and electron jump from valence band into induction band to become free electron (UV-Vis analysis) leads to the occurrence of



transversal and longitudinal mode (Raman spectroscopy) and electron free radicals on the thin film surface. The intensity difference of the free radicals shown in Raman spectroscopy (Figure 4) is caused by In^{3+} doping and radius in the $LiTaO_3$ thin films. For both undoped and In_2O_3 doped $LiTaO_3$ thin films, the experiment results of active phonon frequencies are in good agreement with the literature [12].

4. CONCLUSION

The formation of In_2O_3 doped $LiTaO_3$ thin films with CSD method was succeeded. Analysis of the thin films was conducted using 18D, UV-Vis spectrometer and Raman spectroscopy. The XRD patterns show that the structural properties of $LiTaO_3$ thin film are affected by the indium doping, where the increase in doping concentration causes an increase of crystallite size and lattice parameter values. The reflectance spectrum showed that the films have low reflectance in the ultra-violet range, with the refractive index between 1.703643 and 1.70364377 for undoped film and 6 wt.% In_2O_3 doped film, respectively. The Tauc plot results show that increasing the doping concentration also increases the film band gap energy. The Raman results show that the most significant deviation is 5 cm⁻¹ (A_1TO_4) for the 4 wt.% In_2O_3 doped film. Based on the characterization result, it can be concluded that the $LiTaO_3$ thin films are very promising for application as a light sensor, especially in the ultraviolet region, where 6 wt.% In_2O_3 doped film shows best figure of merit.

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