Aqueous Solution-Grown Crystalline Phosphorus Doped Indium Oxide for Thin-Film Transistors Applications

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Abstract: Solution-grown indium oxide (In2O3) based thin-film transistors (TFTs) hold good prospects for emerging advanced electronics due to their excellent mobility, prominent transparency, and possibility of low-cost and scalable manufacturing; however, pristine In2O3 TFTs suffer from poor switching characteristics due to intrinsic oxygen-vacancy-related defects and require external doping. According to Shanmugam’s theory, among potential dopants, phosphorus (P) has a large dopant–oxygen bonding strength (EM-O) and high Lewis acid strength (L) that would suppress oxygen-vacancy related defects and mitigate dopant-induced carrier scattering; however, P-doped In2O3 (IPO) TFTs have not yet been demonstrated. Here, we report aqueous solution-grown crystalline IPO TFTs for the first time. It is suggested that the incorporation of P could effectively inhibit oxygen-vacancy-related defects while maintaining high mobility. This work experimentally demonstrates that dopant with high EM-O and L is promising for emerging oxide TFTs.

Keywords: thin-film transistors; solution-grown; crystalline oxide semiconductors; phosphorus doped indium oxide

1. Introduction

Transparent oxide semiconductors have emerged as promising materials for a wide range of applications, including thin-film transistors (TFTs) [1–3]; besides, oxide semiconductors could be solution-processed in an ambient atmosphere, showing great potential for scalable manufacturing [4–8]. In2O3 is a representative transparent oxide semiconductor with high mobility due to the large overlap of In 5s orbital; however, the low binding energy of In-O (320.1 KJ/mol) favors oxygen-vacancy formation, resulting in high carrier density [9]. This often leads to high off-state-current and poor subthreshold swing when used as channel material for TFT applications. Previous investigations suggest that doping/alloying is an effective approach to solve this issue [10–17].

Based on Shanmugam’s theory, a promising dopant should have a large dopant–oxygen bonding strength (EM-O) and high Lewis acid strength (L) [11,12]. The large EM-O is able to suppress oxygen-vacancy related defects; however, the introduction of extrinsic dopant would act as scattering centers, causing a drop in mobility [12]. The L can be described as: L = Z/r^2 − 7.7χ_x + 8.0, where Z represents the atomic nucleus charge number, r is the ionic radius, and χ_x designates the element electronegativity [12]. The high L could polarize electron charge away from the oxygen 2p valence band, weakening its activity as scattering centers and, hence, maintaining high mobility [11,12]; therefore, large EM-O and high L are essential for the dopant element. Table 1 tabulated the EM-O...
and L of common dopants for In$_2$O$_3$ host metal oxides [12]. Of all the potential dopants, phosphorus (P) has a large $E_{M-O}$ (599.1 KJ/mol) and high L (10.082), showing great potential for controlling oxygen-vacancy related defects and alleviating dopant impurity carrier scattering. Unfortunately, P-doped In$_2$O$_3$ (IPO) TFTs have not yet been demonstrated and, hence, are worth investigating.

Table 1. Metal-oxygen bonding strength ($E_{M-O}$) and Lewis acid strength (L) of elements.

| Elements | Metal-Oxygen Bonding Strength ($E_{M-O}$, KJ/mol) | Lewis Acid Strength (L) |
|----------|--------------------------------------------------|-------------------------|
| In$^{3+}$ | 320.1                                            | 1.026                   |
| Ga$^{3+}$ | 353.5                                            | 1.167                   |
| Ba$^{2+}$ | 502.9                                            | 1.163                   |
| Mg$^{2+}$ | 363.2                                            | 1.402                   |
| Al$^{3+}$ | 511.0                                            | 3.042                   |
| Sb$^{5+}$ | 434.3                                            | 3.559                   |
| P$^{5+}$  | 599.1                                            | 10.082                  |

In this contribution, we first report the use of P as an efficient dopant for In$_2$O$_3$ TFTs. The IPO channel is fabricated via an aqueous solution route, which enables potential low-cost and expedient manufacturing. The impact of P incorporation is thoroughly investigated. The results show that the incorporation of P could effectively inhibit oxygen-vacancy-related defects while maintaining good mobility. The optimized IPO device (6% P) shows a decent mobility of 6.33 cm$^2$/V s, threshold voltage of 2.97 V, subthreshold swing of 0.60 V/dec, and an excellent $I_{ON}/I_{OFF}$ of $\sim 10^8$, respectively.

2. Results and Discussion

We performed grazing incidence XRD diffraction analysis on the crystalline structure of IPO films with different P proportions. As shown in Figure 1, the broad peaks at $\sim 22.4^\circ$ are attributed to the substrate. The characteristic diffraction peaks of (222) (dominant peak), (400), (440), and (622) are assignable to bixbyite In$_2$O$_3$ (JCPDS 06-0416). No P$_2$O$_5$ or other phases are observed from the diffraction pattern, which means that P addition does not break the In$_2$O$_3$ matrix crystalline structure [18,19]. We suppose that the added P replaces the In sites and maintains the cubic In$_2$O$_3$ structure when the P doping amount is relatively low; however, the diffraction peak intensity gradually decreases, suggesting the reduction of crystallinity after P incorporation [20]. This is because the P element would enhance the local distortions of the InO polyhedral and suppress the number of coordinate In atoms [13,20]. Previous studies also suggest that the addition of different-sized metal cations into the In$_2$O$_3$ could destroy the In$_2$O$_3$ lattice and reduce the crystallinity [10,11,13,20].

AFM measurements shown in Figure 2 indicate that the series of IPO films are highly smooth with RMS roughness of 0.27, 0.24, 0.31, and 0.26 nm for P ratio of 0, 3, 6, and 9%, respectively. The results suggest the absence of prominent grain boundaries in the crystalline IPO films and that P doping has little impact on the surface topography. It should be mentioned that the atomic smooth surface of the channel material is essential for transistor applications.
Figure 1. Grazing incidence XRD patterns for IPO films with indicated P ratio.

Figure 2. AFM images of the surface topography of IPO films with indicated P ratio of (a) 0%, (b) 3%, (c) 6%, and (d) 9%.

Figure 3a represents the optical transmittance spectra of IPO films with a range of P-doping contents. All the IPO films demonstrate excellent transparency (>79% at a wavelength of 550 nm). By using Tauc equation, the optical bandgaps of IPO films (Figure 3b) are calculated to be 3.61 eV (0% P), 3.66 eV (3% P), 3.88 eV (6% P), and 4.19 eV (9% P), respectively. The increased bandgap with P addition could be due to the fact that P can
inhibit the formation of oxygen-vacancy defects in \( \text{In}_2\text{O}_3 \) \[19\]. The increased bandgap is more in line with the larger binding strength of P-O (599.1 KJ/mol) than that of In-O bonding (320.1 KJ/mol), indicating the effective oxygen-vacancy-related defects suppression \[21\].

Figure 3. (a) Optical transmittance spectra of IPO films with various P ratios, and (b) corresponding optical bandgaps derived from the Tauc plots.

XPS is supplemented for further verification. The corresponding P 2p spectra were demonstrated in Figure 4a. Previous reports have summarized the P 2p peak energies \[22\]. The peaks at ~135.6 eV, 133.5 eV, 128.6 eV, and 129.9 eV correspond to \( \text{P}_2\text{O}_5 \), phosphate, phosphide, and elemental P, respectively \[22\]; therefore, in the present work, the characteristic peak at ~133.5 eV is assigned to phosphate (\( \text{PO}_4^{3-} \)). In addition, as shown in Figure 4b, the P/In+ P ratio in the thin film steadily increases from 0 to 3.1, 5.6, and 8.4%, as the P ratio in the precursor increases from 0 to 3%, 6%, and 9%, respectively.

Figure 4. XPS spectra for the IPO films with different P-doping contents. (a) P 2p. spectra, and (b) P-doping ratio between solution and film.

Figures 5 and 6 show the representative transfer and output curves of the IPO TFTs with different P ratios, with electrical parameters summarized in Table 2. The pristine \( \text{In}_2\text{O}_3 \) TFT shows a good mobility of 16.2 cm\(^2\)/V s, but high off-state current and poor subthreshold
slope, which is typical behavior for an undoped-In$_2$O$_3$ device. The P incorporation leads to the great improvement of off-state-current and subthreshold swing, as well as positive shift of threshold voltage; besides, the trap density (including interface and bulk region, see Table 2) also improves after P incorporation, indicating that P doping weakens carrier scattering in IPO [23]. This is because the addition of P could inhibit the oxygen-vacancy-related defects due to the larger $E_{\text{M-O}}$.

Figure 5. Transfer characteristics measured for IPO TFTs with different P ratios.

Figure 6. Output characteristics measured for IPO TFTs with P ratios of (a) 0%, (b) 3%, (c) 6%, and (d) 9%.
Table 2. Summary of electrical properties of IPO TFTs investigated.

| P Ratio (%) | Mobility (cm$^2$/V s) | On/Off Ratio | Threshold Voltage (V) | Subthreshold Swing (V/dec) | Trap Densities (cm$^{-2}$ eV$^{-1}$) |
|-------------|-----------------------|--------------|------------------------|-----------------------------|----------------------------------|
| 0           | 16.2                  | $3.8 \times 10^3$ | -0.57                  | 2.52                        | $8.6 \times 10^{12}$             |
| 3           | 10.8                  | $3.1 \times 10^5$ | 1.98                   | 0.97                        | $2.9 \times 10^{12}$             |
| 6           | 6.33                  | $1.1 \times 10^8$ | 2.97                   | 0.60                        | $1.4 \times 10^{12}$             |
| 9           | 3.52                  | $6.4 \times 10^7$ | 3.27                   | 0.55                        | $1.2 \times 10^{12}$             |

Unfortunately, P introduction would reduce the In 5 s orbital interaction near the conduction band, and, hence, decrease the mobility [10]; however, P has a high Lewis acid strength L, which could polarize electron charge away from the oxygen 2p valence band, weakening carrier scattering centers and, hence, relatively high mobility could be maintained [11,12]. The overall best IPO device (6% P) demonstrates a $\mu$ of 6.33 cm$^2$/V s, $I_{on}/I_{off}$ of $1.1 \times 10^8$, $S$ of 0.60 V/dec, and $V_{th}$ of 2.97 V, respectively, which is sufficient for transistor applications. The mobility of 6.33 cm$^2$/V s is relatively high since mobility larger than 5 cm$^2$/V s is sufficient for liquid crystal display (LCD) applications (the mobility of commercial a-Si is 1 cm$^2$/V s). Although P doping reduces the mobility of pristine In$_2$O$_3$, the other device parameters, especially on/off ratio, is greatly improved.

As shown in Table 3, we have provided the comparison with reported IGZO-related devices in the literature. Our newly designed IPO TFTs are among some of the best results. The newly designed IPO TFTs can be used in flat-panel displays and non-display applications, such as biosensors, photodetectors, memory, and neuromorphic devices.

Table 3. Recent advances of solution-processed IGZO-related TFTs.

| Channel Material | $\mu$ (cm$^2$/V s) | $I_{on}/I_{off}$ | Ref |
|------------------|---------------------|------------------|-----|
| IZO              | 4                   | $10^7$           | [24]|
| IGZO             | 7.5                 | $10^7$           | [25]|
| IGZO             | 8.05                | $10^7$           | [26]|
| IZO              | 2.43                | $10^7$           | [27]|
| IGZO             | 1.38                | $10^5$           | [28]|
| IGZO             | 2.64                | $10^8$           | [29]|
| IPO              | 6.33                | $10^8$           | This work |

3. Experimental Section

Indium nitrate and phosphoric acid were dissolved in DI water with total concentration of 0.2 M and stirred for 2 h before deposition. The desired P/(In + P) atomic contents of 0, 3, 6, and 9% were produced. The IPO layers (~6 nm) were one-step spun at 4500 r.p.m. for 30 s on Si/SiO$_2$ substrates and, then, thermally annealed at 300 °C for 120 min in ambient air. IPO TFTs (W/L = 1500/100 µm) fabrications were completed by evaporating Al source and drain electrodes. The IPO TFT device fabrication process is shown in Figure 7.

Grazing incidence XRD was performed on a Rigaku workstation (Austin, TX, USA). The surface topography information of IPO film was measured via AFM using Bruker Dimension Icon (Beijing, China). The chemical bonding states of IPO were analyzed via X-ray photoelectron spectroscopy (XPS, Thermo Microlab 350, Antigo, WI, USA). The transmittance information of the IPO layer was observed by UV-Vis spectrophotometry with PerkinElmer Lambda 950 (Waltham, MA, USA). Electrical current-voltage measurements of the transistors were carried out in air with a Keithley 2614B digital source meter (San Diego, CA, USA).
Figure 7. The IPO TFT device fabrication process.

4. Conclusions

An efficient and tunable aqueous approach is reported to enhance In$_2$O$_3$ TFTs performance via P doping. It is suggested that P doping in In$_2$O$_3$ TFTs could suppress oxygen-vacancy-related defects that cause electron generation and trapping. The optimal IPO TFTs show enhanced switching characteristics. This work experimentally demonstrates that dopant with high dopant-oxygen bonding strength and Lewis acid strength is a promising candidate and opens a new way to fabricate high performance TFTs at low processing temperature.

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