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- (71) Applicant: PHILLIPS 66 COMPANY [US/US]; Intellectual Property - Legal, P.O. Box 4428, Houston, TX 77210 (US).
- (72) Inventors: WORFOLK, Brian; 2900 Sheridan Road, Bartlesville, OK 74006 (US). EGOAGWU, Nneka, Uguru; 5001 Hollington Drive, Owings Mills, MD 21117 (US).
- (74) Agents: YUEN, Richard, Y. et al.; Phillips 66 Company, Intellectual Property - Legal, P.O. Box 4428, Houston, TX 77210 (US).
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(54) Title: FORMATION OF FILMS FOR ORGANIC PHOTOVOLTAICS

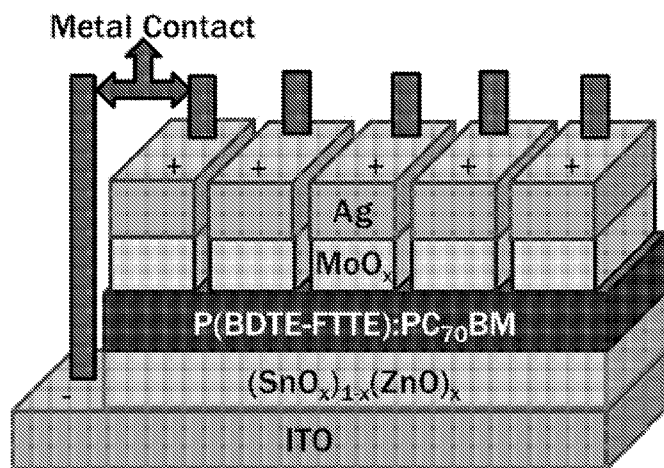


Figure 1

(57) Abstract: A reaction comprising an organic Zn precursor in the amounts of (1-y), an organic Sn precursor in the amounts of y and a base in the amount of (1-y). The product produced from the reaction is  $(\text{SnO}_x)_y\text{ZnO}_{(1-y)}$ .

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TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU,  
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## FORMATION OF FILMS FOR ORGANIC PHOTOVOLTAICS

### CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application is a PCT International application which claims the benefit of and priority to U.S. Provisional Application Ser. No. 62/235,855 filed October 1, 2015 and U.S. Application Serial No. 15/245,676 filed August 24, 2016, entitled "Formation of Films for Organic Photovoltaics," both of which are hereby incorporated by reference.

### STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT

[0002] None.

### FIELD OF THE INVENTION

[0003] This invention relates to the formation of films for organic photovoltaics.

### BACKGROUND OF THE INVENTION

[0004] The present disclosure generally relates to organic solar cells and similar electronic devices. Today's increasing demand for renewable energy resources, especially solar power, is driving researchers to develop low cost, efficient photovoltaic devices. Organic photovoltaics (OPVs) are an attractive route toward solving the terawatt energy problem.

[0005] Solution processed organic photovoltaics have the potential to become a low-cost photovoltaic technology. OPVs can be fabricated on flexible substrates in a roll-to-roll process, which may enable photovoltaics to enter entirely new markets. One of the milestones for commercialization of OPVs is improving device efficiencies, which reduces overall cost. One way of improving device efficiency is through utilizing interfacial charge transport layers.

[0006] Interfacial charge transport layers sandwich the photoactive layer and determine the device polarity, help to collect charges, and transport the charges to the electrodes. Materials for these charge transport layers can be transparent, have low resistance and be chemically stable. The electron transport layer collects and transports electrons mainly generated from the acceptor to the cathode. A low work function interface is required to make Ohmic contact with the organic photoactive layer. ZnO is a common n-type semiconductor used as an electron transport layer in OPVs. ZnO can be processed from a sol-gel solution, and when combined with post-

annealing, produces crystalline thin films at low temperatures (150-200°C). Tin oxide ( $\text{SnO}_x$ ) is known to be highly transparent and conductive but higher post-annealing at 350-500°C is often required. In addition,  $\text{SnO}_x$  has one of the lowest oxygen and water vapor transmission rates for metal oxides, which may aid in organic electronic device stability.

[0007] The low work function of ZnO and high conductivity of  $\text{SnO}_x$  have been previously combined to form mixed metal oxides whereby the ratio of Zn:Sn affects the electronic and physical properties of composites. These zinc tin oxides have application in a number of areas including transistors, sensors, transparent conductors and organic photovoltaics.

[0008] There exists a need for a new low temperature sol-gel solution processing technique for preparing zinc tin oxides with tunable composition.

#### BRIEF SUMMARY OF THE DISCLOSURE

[0009] A reaction comprising an organic Zn precursor in the amounts of (1-y), an organic Sn precursor in the amounts of y and a base in the amount of (1-y) to 1. The product produced from the reaction is  $(\text{SnO}_x)_y\text{ZnO}_{(1-y)}$ .

[0010] A reaction comprising an organic Zn precursor comprising  $\text{Zn}(\text{CH}_3\text{CO}_2)_2$  in the amounts of (1-y), an organic Sn precursor comprising  $\text{Sn}(\text{CH}_3\text{CO}_2)_2$  in the amounts of y, and a base comprising alkanolamine in the amount of (1-y) to 1. The reaction occurs at a temperature less than 225°C to produce  $(\text{SnO}_x)_y\text{ZnO}_{(1-y)}$ .

#### BRIEF DESCRIPTION OF THE DRAWINGS

[0011] A more complete understanding of the present invention and benefits thereof may be acquired by referring to the follow description taken in conjunction with the accompanying drawings in which:

[0012] Figure 1 depicts an inverted device architecture.

[0013] Figure 2 depicts the effect of  $\text{SnO}_x$  content in  $\text{SnO}_x$ :ZnO mixed metal oxide electron transport layers.

[0014] Figure 3 depicts the transmittance of  $\text{SnO}_x$ , 15%  $\text{SnO}_x$  and ZnO films on glass substrates.

[0015] Figure 4 depicts the effect of annealing temperature on the power conversion efficiency of OPV devices with the following architecture: ITO/ $(\text{SnO}_x)_{0.15}:(\text{ZnO})_{0.85}$ /P(BDTE-FTTE)/MoOx/Ag.

[0016] Figure 5 depicts the UPS spectra of ITO, SnO<sub>x</sub>, ZnO and the mixed metal oxide ZTOs.

[0017] Figure 6 depicts the energy diagram illustrating the tunable work function of SnO<sub>x</sub>:ZnO mixed metal oxide composites in alignment with the other layers in the OPV device stack

### DETAILED DESCRIPTION

[0018] Turning now to the detailed description of the preferred arrangement or arrangements of the present invention, it should be understood that the inventive features and concepts may be manifested in other arrangements and that the scope of the invention is not limited to the embodiments described or illustrated. The scope of the invention is intended only to be limited by the scope of the claims that follow.

[0019] The present embodiment discloses a reaction of an organic Zn precursor in the amounts of (1-y), an organic Sn precursor in the amounts of y; and a base in the amount of (1-y) to 1. The resultant product is (SnO<sub>x</sub>)<sub>y</sub>ZnO<sub>(1-y)</sub>.

[0020] In one embodiment the organic zinc precursor comprises Zn(CH<sub>3</sub>CO<sub>2</sub>)<sub>2</sub> or Zn(CH<sub>3</sub>CO<sub>2</sub>)<sub>2</sub>\*2H<sub>2</sub>O.

[0021] In one embodiment the organic tin precursor comprises Sn(CH<sub>3</sub>CO<sub>2</sub>)<sub>2</sub>.

[0022] In another embodiment the base is an alcohol. Examples of bases that can be used including amines or alkanolamines.

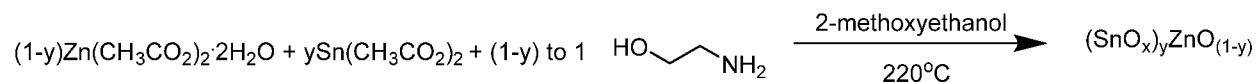
[0023] In yet another embodiment, the reaction also comprises a solvent. The solvent can be used to dissolve either the zinc precursor or the tin precursor. One example of a solvent that can be used is water, alcohol, aminoalcohol, carboxylic acid, glycol, hydroxyester, aminoester or a mixture. Some examples include: 2-methoxyethanol, methanol, ethanol, propanol, butanol, pentanol, hexanol, ethylene glycol, ethoxyethanol, methoxyethanol, ethoxypropanol, ethoxyethanol, dimethoxyglycol, N,N-dimethylformamide.

[0024] In one embodiment, (SnO<sub>x</sub>)<sub>y</sub>ZnO<sub>(1-y)</sub> is used as an electron transport layer for an organic photovoltaic device. In another embodiment the organic photovoltaic device has an inverted device architecture. An inverted device architecture has the positive and negative electrodes reversed. Figure 1 depicts an inverted device architecture which employs indium tin oxide as the cathode and silver as the anode. In this type of device, it is theorized that electrons need to move from the polymer:fullerene active layer to the cathode. Electrons are transported

from the photoactive layer by the electron transport layer, and extracted to the transparent cathode.

**[0025]** In one embodiment,  $(\text{SnO}_x)_y\text{ZnO}_{(1-y)}$  is a sol-gel solution.

**[0026]** In another embodiment,  $(\text{SnO}_x)_y\text{ZnO}_{(1-y)}$  was prepared by dissolving zinc acetate dihydrate or tin(II) acetate in 2-methoxyethanol and ethanolamine. One example of the reaction is shown below:



**[0027]** Examples:

**[0028]** Solution formation

**[0029]** A ZnO sol-gel solution was prepared by mixing 0.33 g  $\text{Zn}(\text{CH}_3\text{CO}_2)_2$  in 3 mL of 2-methoxyethanol with 92  $\mu\text{L}$  of ethanolamine. Similarly SnOx sol-gel solutions were prepared by dissolving 0.36 g of  $\text{Sn}(\text{CH}_3\text{CO}_2)_2$  in 3.5 mL of 2-methoxyethanol, and 99  $\mu\text{L}$  of ethanolamine. ZnO & SnOx were studied independently and as a mixed metal oxide system. Mixed sol-gel solutions were prepared from stock zinc and tin solutions. The amount of Sn in the mixed solution could be (5, 10, 15, 70, 95) vol%. In this embodiment the solutions were stirred for at least an hour before spin casting on indium tin oxide.

**[0030]** Device Fabrication

**[0031]** The photoactive layer consisted of the donor polymer poly(4,8-bis(5-2-ethylhexyl)thiophene-2-yl)benzo[1,2-b:4,5-b']dithiophene-2-ethylhexyl-4,6-dibromo-3-fluorothieno[3,4-b]thiophene-2-carboxylate (P(BDTE-FTTE)) and acceptor [6,6]-phenyl-C70-butyric acid methyl ester (PCBM) at a ratio of 1:1.6, respectively. The total solution concentration was 26 mg/mL in o-xylene. The photoactive layer solution was stirred and heated at 80 °C overnight in a nitrogen filled glove box. The next day 2.5 vol% of 1,8-diiodooctane was added and the solution was heated on the hot plate at 80 °C for an hour. The solution was then filtered with a 2.7  $\mu\text{m}$  glass fiber syringe filter.

**[0032]** Indium tin oxide patterned glass substrates were cleaned by successive ultrasonifications in detergent, deionized water, acetone, and isopropanol. Each 15 min step was repeated twice and the freshly cleaned substrates were left to dry overnight at 80 °C. Preceding fabrication, the substrates were further cleaned for 30 min in a UV-ozone chamber and the electron transport layer was immediately spin coated on top.

**[0033]** Single component or mixed metal oxide solutions were filtered directly onto indium tin oxide with a 0.25  $\mu\text{m}$  poly(tetrafluoroethylene) filter and spin cast at 5000 rpm for 40 s. Film were then annealed at 220  $^{\circ}\text{C}$  for 15 min, and directly transferred into a nitrogen filled glove box. ZnO films were annealed at 170  $^{\circ}\text{C}$ .

**[0034]** The photoactive layer was deposited on the electron transport layer via spin coating at 1200 rpm for 40 s and directly transferred into a glass petri dish to solvent anneal for 1 h. After solvent annealing, the substrates were loaded into the vacuum evaporator where MoO<sub>x</sub> (hole transport layer) and Ag (anode) were sequentially deposited by thermal evaporation. Deposition occurred at a pressure of  $1 \times 10^{-6}$  torr. MoO<sub>x</sub> and Ag had a thickness between 10 nm and 200 nm, respectively. Samples were then were then encapsulated with glass using an epoxy binder and treated with UV light for 3 min.

**[0035]** Table 1 depicts the photovoltaic parameters of ZnO and SnO<sub>x</sub> electron transport layer with the following device architecture: ITO/ETL/P(BDTE-FTTE):PCBM/MoO<sub>x</sub>/Ag.

ETL	J <sub>sc</sub> (mA/cm <sup>2</sup> )	V <sub>oc</sub> (V)	FF (%)	PCE (%)	R <sub>s</sub> ( $\Omega$ cm <sup>2</sup> )	R <sub>sh</sub> ( $\Omega$ cm <sup>2</sup> )	Work Function (eV)
ZnO	15.1	0.774	61.6	7.21	9.88	816	3.75
(SnO <sub>x</sub> ) <sub>0.05</sub> (ZnO) <sub>0.95</sub>	14.8	0.760	55.2	6.67	5.06	288	3.68
(SnO <sub>x</sub> ) <sub>0.15</sub> (ZnO) <sub>0.85</sub>	16.0	0.779	66.9	8.28	5.64	832	3.74
(SnO <sub>x</sub> ) <sub>0.75</sub> (ZnO) <sub>0.25</sub>	15.6	0.713	55.8	6.17	12.4	623	3.93
(SnO <sub>x</sub> ) <sub>0.95</sub> (ZnO) <sub>0.05</sub>	15.8	0.737	61.3	7.08	15.4	476	4.13
SnO <sub>x</sub>	15.7	0.757	62.3	7.41	6.87	769	4.15

Table 1

**[0036]** Using ZnO as the electron transport layer resulted in an average power conversion efficiency (PCE) of 7.21%, compared to the average power conversion efficiency of SnO<sub>x</sub> of 7.41%. The tin oxide ETLs had higher short-circuit current density (J<sub>sc</sub>) and lower series resistance (R<sub>s</sub>) which can be attributed to its superior transparency and conductivity properties, respectively. ZnO devices had higher open-circuit voltages (V<sub>oc</sub>) presumably due to better interfacial energy alignment with the photoactive layer as a result of its lower bulk work function.

**[0037]** Mixed Metal Oxide Electron Transport Layers

**[0038]** In order to determine whether there is any effect of combining low work function ZnO with conductive SnO<sub>x</sub>, a range of mixed metal oxide compositions were prepared, where the SnO<sub>x</sub> component ranged from 5% to 95% (by volume). As the SnO<sub>x</sub> content increased, there

is a peak in photovoltaic performance at 15% SnOx. On either side of 15% SnOx the performance drops significantly as seen in Figure 2.

**[0039]** The high conductivity of tin oxide and the high transparency of ZnO have been combined at an optimal ratio of (SnOx)<sub>0.15</sub>(ZnO)<sub>0.85</sub>, which resulted in an average PCE of 8.28%. This is significantly higher than the photovoltaic performance of the pure ZnO or SnOx thin films. This composition had the best performance in all photovoltaic parameters except for the Rs. In general, ZTOs with a higher SnOx content had a superior Jsc, likely the result of higher transparency, but had a lower Voc due to the higher work functions of SnOx rich composites, as reported in the literature.

**[0040]** To further optimize the processing conditions for 15% SnOx, the films were annealed at 170, 200 and 220 °C to elucidate the effect on the photovoltaic performance. Table II depicts photovoltaic parameters of different annealing temperatures on (SnOx)<sub>0.15</sub>:(ZnO)<sub>0.85</sub> electron transport layer organic photovoltaic devices.

Annealing Temperature (°C)	Jsc (mA/cm <sup>2</sup> )	Voc (V)	FF (%)	PCE (%)	Rs (Ω cm <sup>2</sup> )	Rsh (Ω cm <sup>2</sup> )
170	15.8	0.801	61.1	7.72	5.39	624
200	16.2	0.804	63.6	8.29	5.35	759
220	16.6	0.803	63.4	8.45	5.30	754

Table II

**[0041]** At 170 °C, the average PCE of devices was 7.72%. By increasing the ETL annealing temperature to 200 °C and 220 °C, the PCE increased to 8.29% and 8.45%, respectively. A major contributor to the increase in PV performance was the increase in the Jsc and FF. At lower annealing temperatures, the ZTO composite likely has poor crystallinity, which improves with higher annealing temperatures. However, by annealing the electron transport layer at 220 °C, device efficiencies up to 8.99% were attained. The sol-gel synthesis for ZTO thin films is able to obtain high performance at significantly lower annealing temperatures compared to the standard ZnO films. Annealing the electron transport layer at lower temperatures is beneficial when transferring processing to flexible plastic substrates and roll-to-roll processing.

**[0042]** Optical Properties of Zinc Tin Oxide Films

**[0043]** As light must first pass through the electron transport layer to the photoactive layer to generate charges, high transparency of the film across the solar spectrum is critical. The transparency of SnOx, ZnO, and 15% SnOx was characterized on glass substrates and the



transmittance spectra are presented in Figure 3. It is common for the scientific community to benchmark the transmittance of transparent conductors at 550 nm. From Figure 3, the transparency of the 15% SnOx sample is superior to the single component metal oxides, reaching 98.8% transparency at 550 nm. ZnO and SnOx films are 96.9% and 95.2% transparency at 550 nm. The superior optical properties of 15% SnOx are reflected in this composite obtaining the highest average Jsc in OPV devices. Allowing more photons to pass through the cathode and electron transport layer can increase the absorption of the photoactive layer, resulting in higher Jsc.

**[0044]** The ZnO film shows an excitonic peak at ~346 nm, which is characteristic of small ZnO crystallites. When adding 15% SnOx to ZnO, the peak shifts to higher energy at ~325 nm. This blue-shift is characteristic with a reduction in the crystallite size of ZnO by the addition of SnOx. The peak width is significantly wider, indicating a higher degree of polydispersity of ZnO crystallite sizes in these films. The SnOx spectrum is nearly featureless with a very small electronic transition at ~ 475 nm. As this peak intensity is very small, the SnOx film has a low degree of crystallization. As shown in Figure 4, higher annealing temperatures may increase the crystallinity of SnOx and mixed ZTO films, however higher temperatures must be balanced by processing cost and adaptability to flexible plastic substrates.

**[0045]** Work Function of Mixed Metal Oxide Films

**[0046]** To understand the role of SnOx in the mixed metal oxide films we determined the work function of the films using ultraviolet photoelectron spectroscopy (UPS). UPS is analogous to X-ray photoelectron spectroscopy (XPS) but uses ultraviolet radiation instead. Since the power of UV light is lower than X-rays, UPS is even more surface sensitive compared to XPS. As such, UPS typically characterizes the top 1-3 nm surface of films. In photoelectron spectroscopy, the addition of UV energy ( $h\nu$ ) and kinetic energy (KE) of emitted electrons is equivalent to the binding energy (BE) of electrons within a specific atomic orbital. This is formalized into the following equation:

**[0047]** 
$$BE = KE + h\nu \quad (1)$$

**[0048]** UPS detects both photoelectrons and secondary electrons. The cutoff of the secondary electron peak at high binding energy is concomitant with the film's surface work function, which is the minimum amount of energy required to remove an electron from a film to vacuum. The

work function of anodes, cathodes and carrier transport layers is critical in organic photovoltaics as it determines the device's polarity, as well as carrier extraction efficiency.

**[0049]** The UPS spectra of mixed metal oxide films are seen in Figure 5. The spectra are plotted showing the secondary electron cutoff region where the work function is determined. The work function of the ITO cathode is 4.65 eV. In order for ITO to collect electrons, the work function must be lowered to increase electron specificity. Both ZnO and SnO<sub>x</sub> decrease the work function to 3.75 eV and 4.15 eV, respectively. For composite films at 95% and 70% SnO<sub>x</sub>, the work function is in between the single component metal oxide and is 4.13 eV and 3.93 eV. Further decreasing the SnO<sub>x</sub> content reduces the work function lower than ZnO-only films to reach 3.73 eV at 30% SnO<sub>x</sub> and 3.68 eV at 5% SnO<sub>x</sub>. These two compositions also have the lowest R<sub>s</sub> as seen in Table I, which indicates a reduction of resistive losses in the OPV devices. Further reducing the work function beyond that of ZnO is particularly interesting as the mixed metal combination obtains different physical properties compared to the individual materials on their own.

**[0050]** An energy band diagram for the organic photovoltaic device architecture is presented in Figure 6. The figure reiterates that decreasing SnO<sub>x</sub> content in ZTO films reduces the work function. Ideally the work function of the electron transport layer should be less than the lowest unoccupied molecular orbital energy of PCBM. This is the case for 15% and 5% SnO<sub>x</sub> as well as 100% ZnO. For the ZTO composites, lowering the tin content to 15% improves photovoltaic performance as a result of improved interfacial energy alignment. ZTO composites less than 15% tin have a reduction in performance. This may be the result of lowering the film conductivity with increasing zinc content.

**[0051]** ZTO films contain 15-20 atomic% of acetate as characterized with X-ray photoelectron spectroscopy. Table III below depicts the atomic concentration of ZTO films cast from 65% and 35% diluted sol-gel solutions.

	65%				35%	
	Room Temperature	170°C	210°C	240°C	170°C	210°C
O	43.6	43.6	44.3	45.3	43.4	44.3
C-C	17.6	14.9	12.7	10.4	20.4	25.2
COOH	11.3	10.3	8.5	6.3	4.8	3.4

Zn	26.0	28.9	31.4	35.0	27.7	22.4
Sn	0.7	1.4	2.1	2.2	3.8	4.8
N	0.8	0.9	1.1	0.8	-	-

Table III

**[0052]** In closing, it should be noted that the discussion of any reference is not an admission that it is prior art to the present invention, especially any reference that may have a publication date after the priority date of this application. At the same time, each and every claim below is hereby incorporated into this detailed description or specification as an additional embodiment of the present invention.

**[0053]** Although the systems and processes described herein have been described in detail, it should be understood that various changes, substitutions, and alterations can be made without departing from the spirit and scope of the invention as defined by the following claims. Those skilled in the art may be able to study the preferred embodiments and identify other ways to practice the invention that are not exactly as described herein. It is the intent of the inventors that variations and equivalents of the invention are within the scope of the claims while the description, abstract and drawings are not to be used to limit the scope of the invention. The invention is specifically intended to be as broad as the claims below and their equivalents.

## CLAIMS

1. A reaction comprising:
  - an organic Zn precursor in the amounts of (1-y);
  - an organic Sn precursor in the amounts of y; and
  - a base in the amount of (1-y) to 1,to produce  $(\text{SnO}_x)_y\text{ZnO}_{(1-y)}$ .
2. The reaction of claim 1, wherein the organic Zn precursor comprises  $\text{Zn}(\text{CH}_3\text{CO}_2)_2 \cdot 2\text{H}_2\text{O}$ .
3. The reaction of claim 1, wherein the organic Sn precursor comprises  $\text{Sn}(\text{CH}_3\text{CO}_2)_2$ .
4. The reaction of claim 1, wherein the base is an alcohol.
5. The reaction of claim 1, wherein the base is alkanolamine.
6. The reaction of claim 1, wherein the reaction also comprises a solvent.
7. The reaction of claim 1, wherein the solvent is 2-methoxyethanol.
8. The reaction of claim 1, wherein the reaction occurs at a temperature above room temperature.
9. The reaction of claim 1, wherein the reaction occurs at a temperature greater than 150°C.
10. The reaction of claim 1, wherein the reaction occurs at a temperature less than 250°C.
11. The reaction of claim 1, wherein the reaction occurs at a temperature less than 225°C.
12. The reaction of claim 1, wherein  $(\text{SnO}_x)_y\text{ZnO}_{(1-y)}$  contains from about 10 to 25% atomic% of acetate as characterized with x-ray photoelectron spectroscopy.
13. The reaction of claim 1, wherein  $(\text{SnO}_x)_y\text{ZnO}_{(1-y)}$  is used as an electron transport layer for an organic photovoltaic device.
14. A reaction comprising:
  - an organic Zn precursor comprising  $\text{Zn}(\text{CH}_3\text{CO}_2)_2$  in the amounts of (1-y);
  - an organic Sn precursor comprising  $\text{Sn}(\text{CH}_3\text{CO}_2)_2$  in the amounts of y; and
  - a base comprising alkanolamine in the amount of (1-y),wherein the reaction occurs at a temperature less than 225°C to produce  $(\text{SnO}_x)_y\text{ZnO}_{(1-y)}$ .

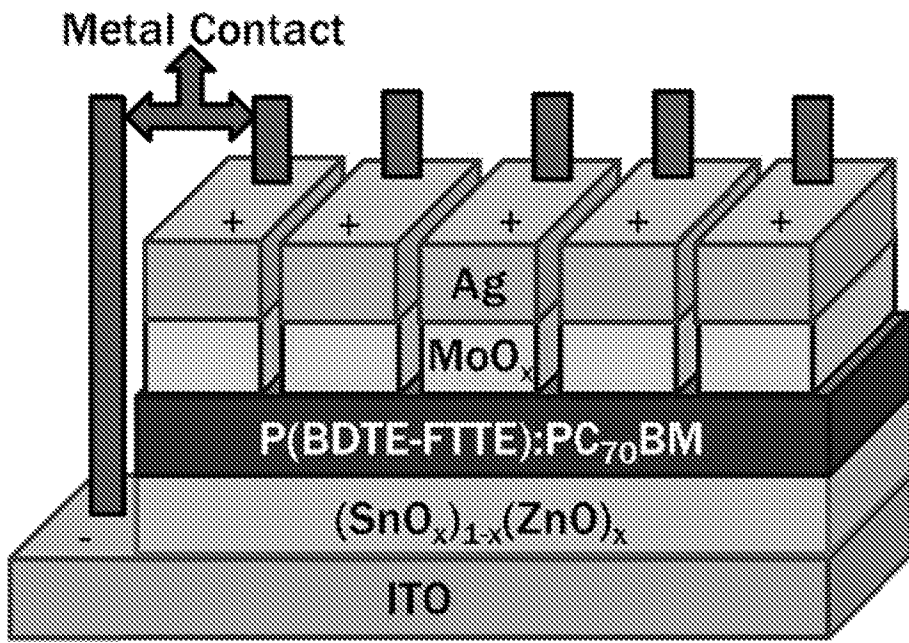


Figure 1

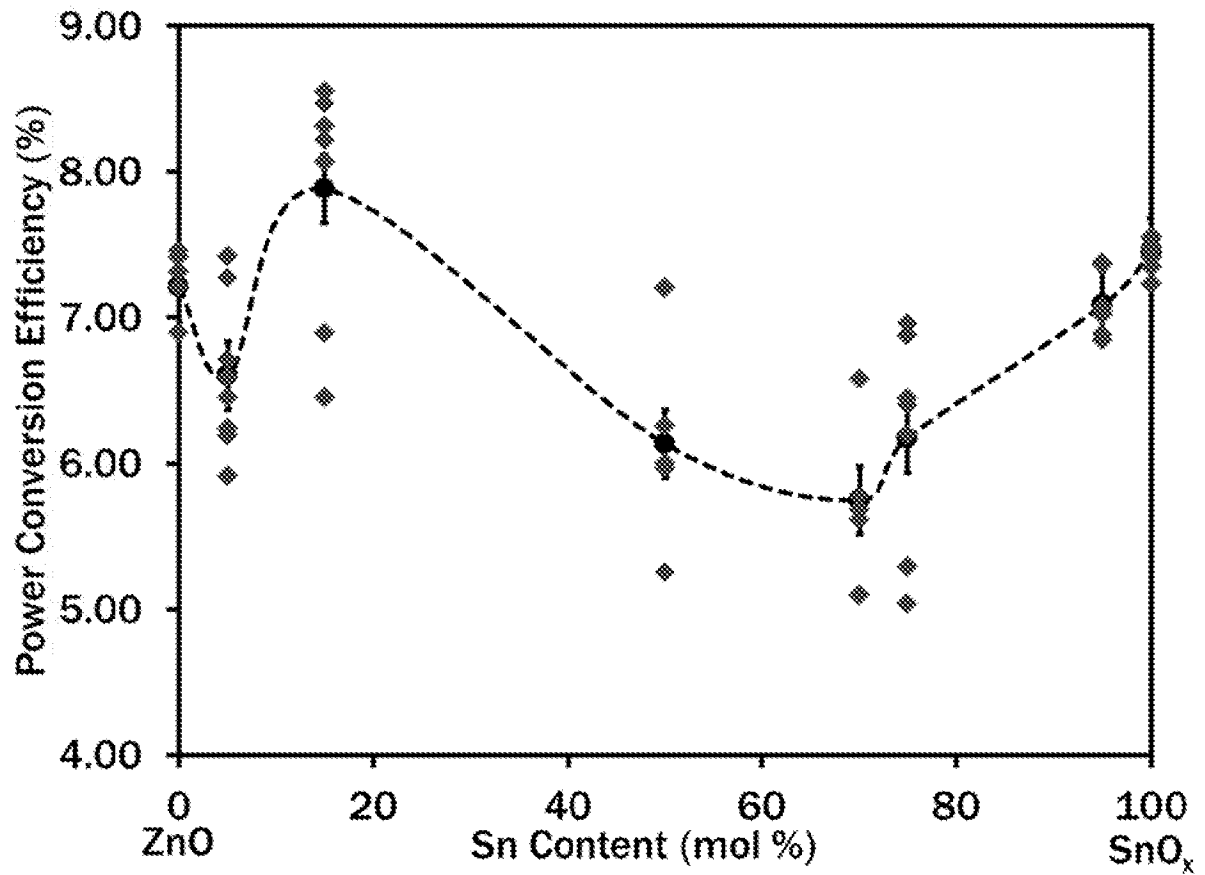


Figure 2

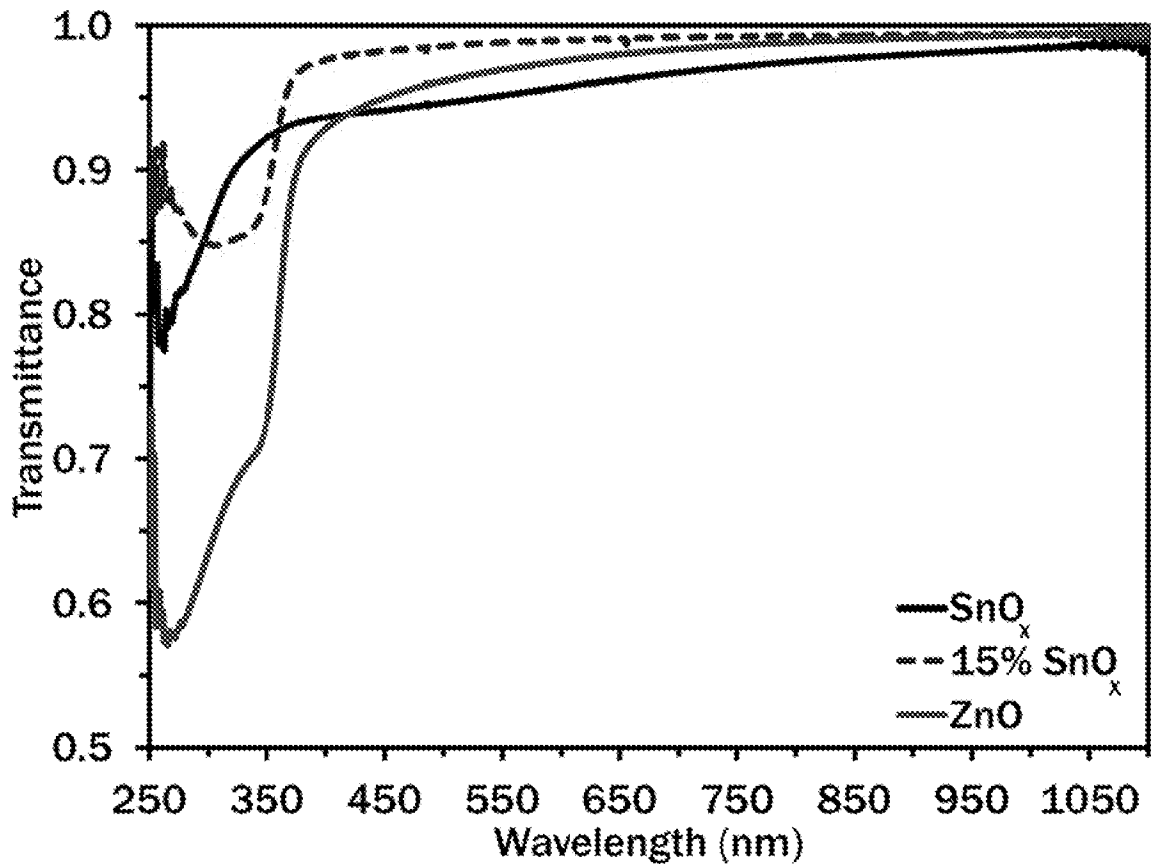


Figure 3

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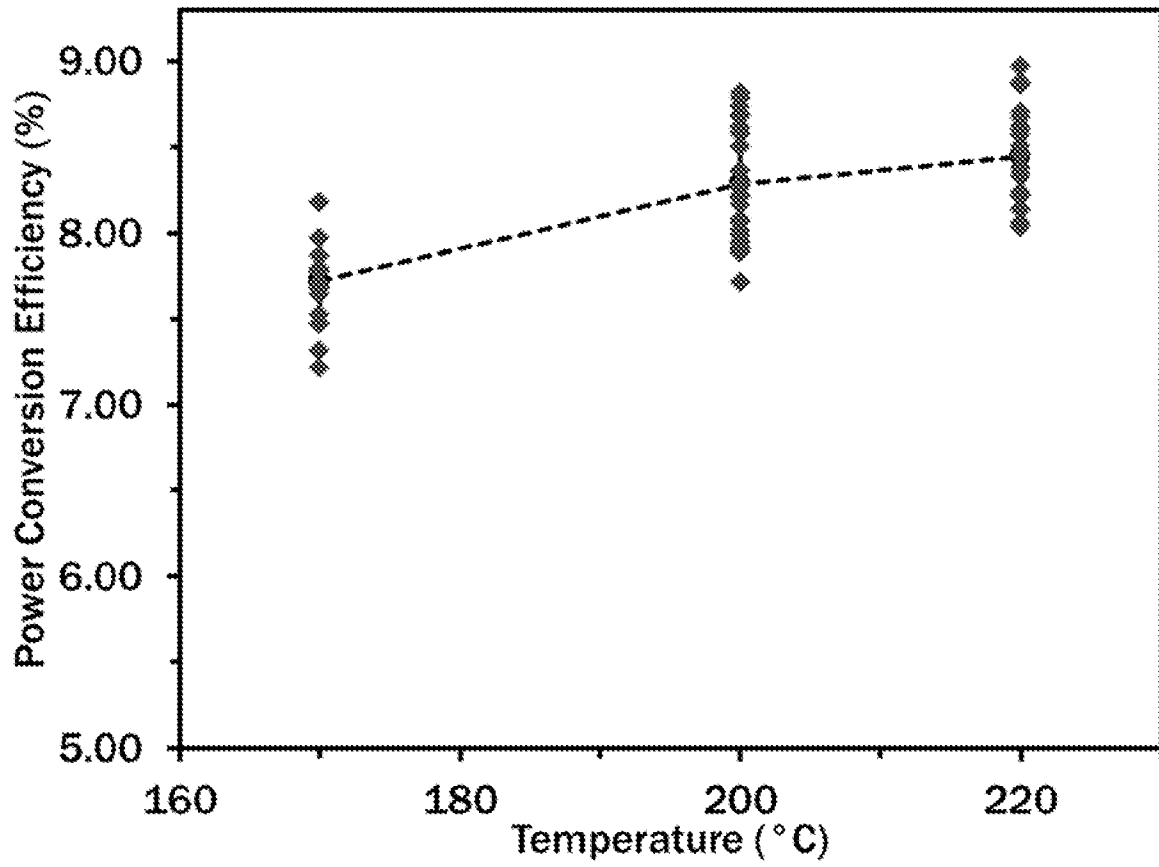


Figure 4



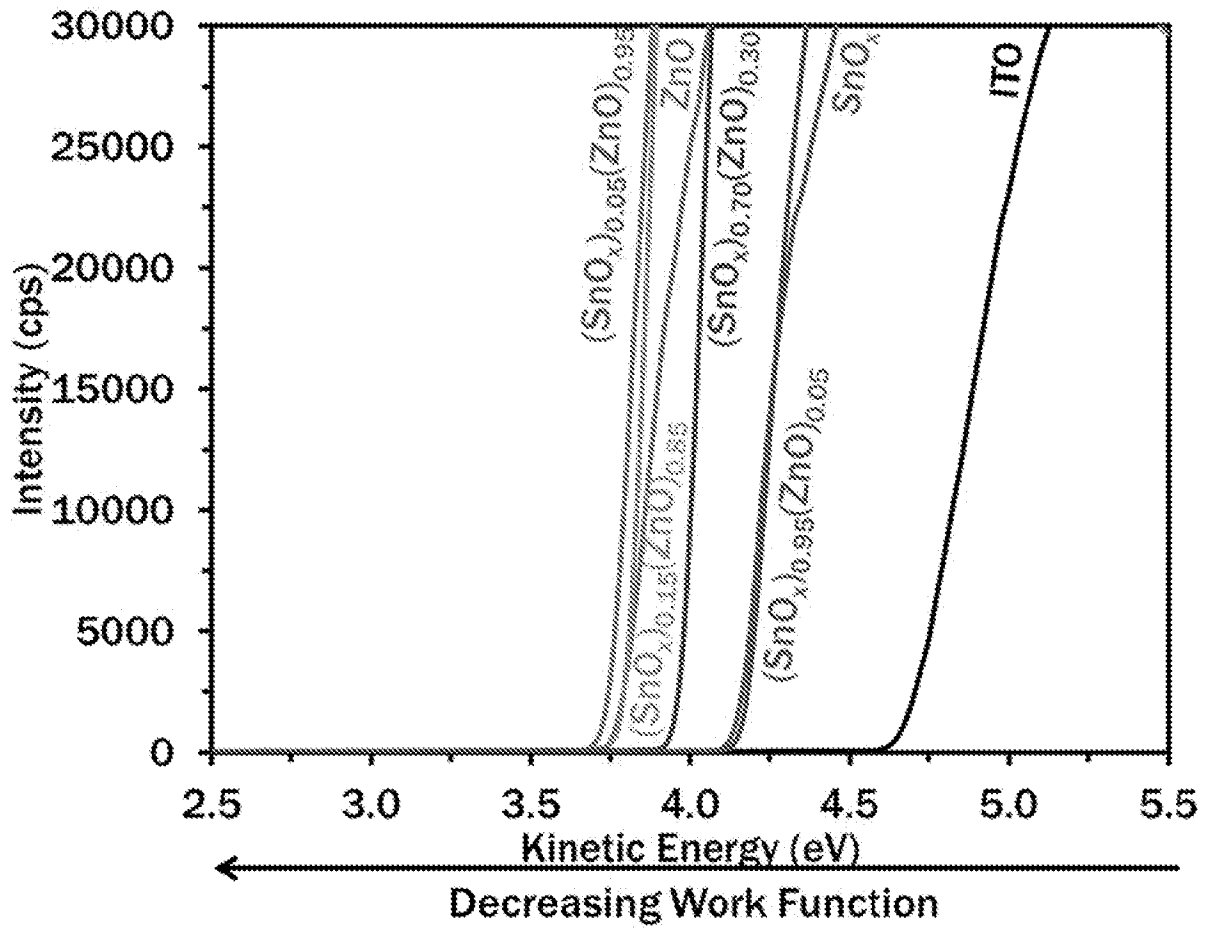


Figure 5

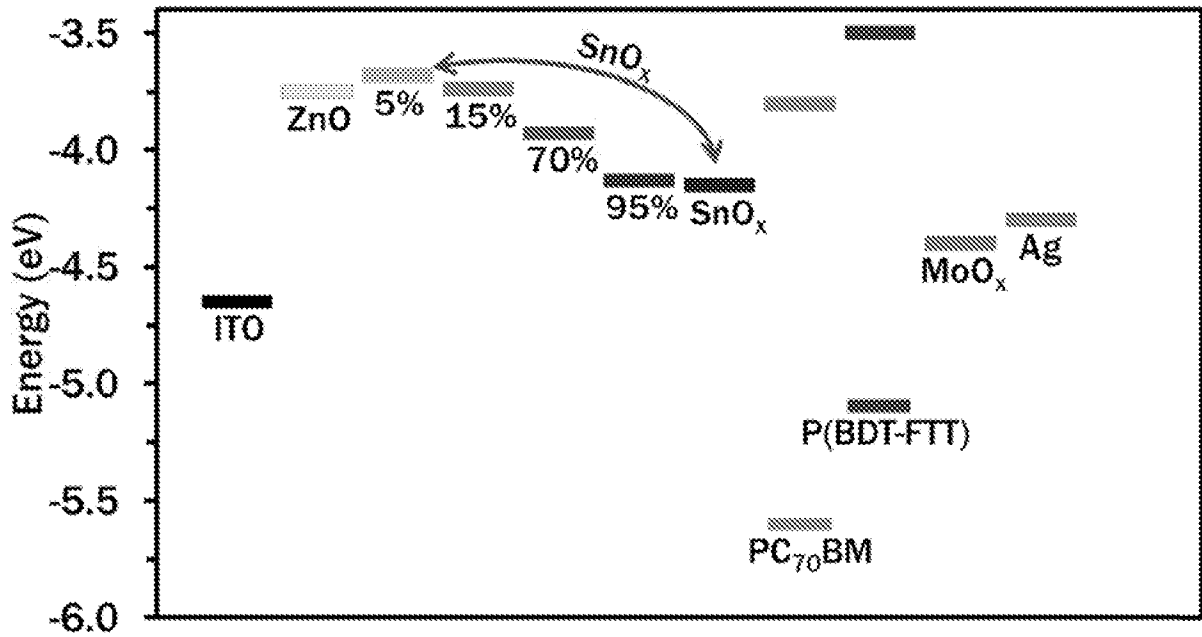


Figure 6

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US16/48389

<b>A. CLASSIFICATION OF SUBJECT MATTER</b> IPC(8) - C23C 6/00, 18/08, 18/12, 30/00; H01L 21/02 (2016.01) CPC - C23C 6/00, 18/08, 18/1216, 18/1254, 30/00; H01L 21/02554 According to International Patent Classification (IPC) or to both national classification and IPC		
<b>B. FIELDS SEARCHED</b> Minimum documentation searched (classification system followed by classification symbols) IPC(8) - C23C 6/00, 18/08, 18/12, 30/00; H01L 21/02 (2016.01) CPC - C23C 6/00, 18/08, 18/1216, 18/1254, 30/00; H01L 21/02554, 21/02565 ; USPC - 257/43, E29.273; 438/478 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) PatSeer (US, EP, WO, JP, DE, GB, CN, FR, KR, ES, AU, IN, CA, INPADOC Data); Google; Google Scholar; EBSCO; IP.com; organic, photovoltaic, tin acetate, Sn, zinc, zinc acetate, Zn, sol-gel, solar cell, organic Zn precursor, organic Sn precursor, base, (SnOx)yZnO (1-y), ZnO, SnOx, Sn(CH3CO2)2, Zn(CH3CO2)2, zinc tin oxide, ZTO, x-ray photoelectron spectroscopy, XPS, acetate, concentration		
<b>C. DOCUMENTS CONSIDERED TO BE RELEVANT</b>		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X ---	US 2010/0251936 A1 (KIM, HJ, et al.) October 7, 2010; paragraphs [0006], [0052]	1-8, 10-11, 14 ---
Y		9, 13
Y	US 2013/0292668 A1 (KOREA INSTITUTE OF SCIENCE AND TECHNOL) November 7, 2013; paragraphs [0015], [0027]	9
Y	WO 2014/191076 A1 (MERCK PATENT GMBH) December 4, 2014; page 15, lines 7-8, 20-26; claim 3	13
A	(KIM, YJ et al.) Impact of the cation composition on the electrical performance of solution-processed zinc tin oxide thin-film transistors. ACS applied materials & interfaces. 2014. vol. 6; abstract; page 14027, column 2, 3rd paragraph	1-14
A	(JEONG, S et al.) Solution-processed zinc tin oxide semiconductor for thin-film transistors. The Journal of Physical Chemistry C. 2008. vol. 112; pages 11082-11085	1-14
<input type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.		
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family		
Date of the actual completion of the international search		Date of mailing of the international search report
8 October 2016 (08.10.2016)		27 OCT 2016
Name and mailing address of the ISA/US Mail Stop PCT, Attn: ISA/US, Commissioner for Patents P.O. Box 1450, Alexandria, Virginia 22313-1450 Facsimile No. 571-273-8300		Authorized officer Shane Thomas PCT Helpdesk: 571-272-4300 PCT OSP: 571-272-7774