

A Review On The Absorber Materials In Dye Sensitized Solar Cell

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Abstract—This paper reports on the fabrication of dye sensitized solar cell using various types of photo electrodes. They have been widely used because of their unique properties including high absorbance coefficient, chemical stability and electrochemical activity at nanoscale. In this work, the growth conditions will be optimized in order to determine the best power conversion efficiency.

Keywords—*thin films; dye sensitized solar cell; band gap; semiconductor*

I. INTRODUCTION

Recently, a number of works which describe the power conversion efficiency of fabricated dye sensitized solar cells (DSSC) have been done by many researchers and successfully published their results in many papers. The investigation of this type of solar cell is important due to many advantages such as inexpensive [1, 2], efficient solar energy conversion [3], flexibility [4] and easy production [5]. Generally, the dye sensitized solar cell consists of a photoelectrode and a catalytic electrode [6] with an electrolyte between them [7]. In order to study the photoelectrochemical properties of nanoparticles, various parameters including conversion efficiency, open circuit voltage, close circuit current density, fill factor, interface charge resistance and an incident photon will be investigated. Researchers understand that these photoelectrochemical properties are strongly depend on the morphological properties of nanoparticles, spectroscopic characteristics of dyes and the electrical behavior of electrolytes.

Nowadays, scientists have to look for renewable energy sources because of the consumption of fossil fuels are increasing day by day [8]. In this work, dye sensitized solar cell will be fabricated by using different photo electrodes. Then, the performance of solar cell will be evaluated based on photoelectrochemical parameters.

II. LITERATURE SURVEY

Rani and Ernest [9] designed the dye sensitized solar cell using different photo electrodes, namely zinc oxide (ZnO) and gallium doped zinc oxide thin films. The experiment results show that gallium doped zinc oxide films have the larger surface area. Therefore, it reaches the efficiency of 4.3% than the zinc oxide electrode of 2.5%. In other words, the conversion efficiency in latter case is low due to the order less

structure of film which was supported by the scanning electron micrographs. On the other hand, they also prepared gallium doped zinc oxide films of different concentrations of gallium. They conclude that the efficiency of the solar cell increases for every increment in the concentration of gallium from 0.0001 M (3.9%), 0.0002 M (4.1%) to 0.0003 M (4.3%).

The influence of zinc oxide coated titanium dioxide (TiO₂) working electrode in dye sensitized solar cell was studied by Chou et al [10]. They presented some results such as the band gap and atomic percentage of zinc dispersed in the zinc oxide coated TiO₂ films were increased at longer soaking time and with a higher concentration of zinc acetate dihydrate. It is due to more zinc oxide was deposited on the TiO₂ films, then might facilitate the creation of the energy barrier in the working electrode of a dye sensitized solar cell. As a result, conversion efficiency of the zinc coated TiO₂ electrode (6.7%) exceeds that of the conventional solar cell with a TiO₂ electrode (5.9%).

Anca et al [11] investigated the influence of surfactant in morphology and optical properties of zinc oxide nanostructures. They found that more uniform distribution of particles dimensions and a higher crystallinity for triton-100 as a surfactant. Also, the mean diameter of nanoparticles about 19 nm and 30 nm could be detected for the films prepared using triton X-100 and PEG 200, respectively. Lastly, they conclude that the higher values of efficiencies (1.01-1.19%) in the case of zinc oxide in the presence of triton X-100 than that of films obtained in the presence of PEG 200 (0.24%).

Wu et al [12] developed dye sensitized solar cell based on nickel oxide (NiO) electrodes. They point out that the conversion efficiency of solar cell based on improved hydrothermal method was twice as high as that of primary hydrothermal method. On the other hand, they reported that the reaction solution concentration has great effect on the properties of the NiO films. For example, the morphology of NiO changes from flocks consisting of nanosheets to flower like microspheres as shown in scanning electron microscopy analysis. Also, conversion efficiency is increased from 0.015% to 0.036% as the solution concentration was increased from 15 to 50 mM in primary NiO photocathodes.

ZnO and Al-doped ZnO thin films were deposited on transparent conducting oxide glass using glass rod

spreading method by Suri et al [13]. They choose ZnO in their experiment due to many reasons such as its stability against photo corrosion and photochemical properties similar to TiO₂. They found that the lower conversion efficiency of the Al doped ZnO (0.6 %) compared to ZnO (1.43 %) because of lower injection efficiency and less porosity in this type of solar cell.

Supphadate & Sasimonton [14] demonstrated an improvement of conversion efficiency of DSSC by addition of stannic oxide (SnO₂) into ZnO photoanode. They proposed that the power conversion efficiency reaching a maximum value of 1.75 % where additive SnO₂ weight ratio of 7 %, indicating that the conversion efficiency changed with SnO₂ weight ratio. They explain that the conversion efficiency is reached by reduction of transfer resistance to induce pathway direction for electron transfer in photoanode.

The cupric oxide (CuO) nanoparticles were conducted by Jitendra et al [15]. The obtained nanoparticles show monoclinic structure and exhibit spherical morphology as indicated in X-ray diffraction (XRD) and scanning electron microscopy (SEM) investigations. Additionally, these compounds show good surface for the electrocatalytic activity towards the reduction of I₃⁻ (triiodide) to I⁻ iodide ions in redox electrolyte as reported in cyclovoltammetry analysis. Lastly, they claim the conversion efficiency of 3.4 % was achieved in their fabricated solar cell by using CuO nanoparticles.

Electrophoretic deposition technique was used for the fabrication of TiO₂ films on the conductive substrate by Masood et al [16]. They analyzed that an increase of deposition cycle lead to linear increases of film thickness and cell efficiency. For example, the power conversion efficiency was increased from 0.074% to 6.56% as the film thickness was increased from 1.5 μm to 24 μm. On the other hand, the mean sizes of nanoparticles were increased as the annealing temperature was increased from 150 °C (21 nm) to 500 °C (26.5 nm) according to the scanning electron microscopy results. Lastly, they suggest that the optimized annealed temperature should be 500 °C because of successfully yielded the highest conversion efficiency of 6.6 %.

Zinc oxide nanoparticles were electrodeposited on TiO₂ photoanode by Mozaffari et al [17] in order to suppress the electron recombination process. In order to achieve this objective, various deposition times and concentrations of zinc should be examined in their experiments. They observe that a large amount of ZnO was covered on TiO₂ nanoparticles at high concentration of zinc (1 x 10⁻³ M) and longer deposition time (300 s). Lastly, they suggest that the best conditions to produce higher conversion efficiency of 4.56 % were at deposition time of 15 seconds and concentration of zinc was 1 X10⁻⁶ M by comparing the results of all accomplished experiments.

Chou et al [18] studied the applicability of n-type TiO₂ and p-type NiO on the fluorine doped tin oxide

substrate in dye sensitized solar cell. They also analyzed the influence of the mass ratio of TiO₂ to Ni and the number of coatings of TiO₂ particles on the conversion efficiency. The power conversion efficiency increases from 3.1 % to 3.6 % as the mass ratio of TiO₂ to Ni changes from 10:0.1 to 10:0.2, indicating that conversion efficiency increases with increasing the mass of Ni in TiO₂/NiO particles.

TABLE I. CHEMICAL BATH DEPOSITED THIN FILMS

| |
|-------------------------------------|
| NiS [20] |
| CdS [21] |
| Cu-Sn-S [22] |
| Cd-Zn-S [23] |
| Cu ₂ S [24] |
| Cu-Cd-S [25] |
| FeS ₂ [26] |
| SnS [27, 28] |
| ZnS [29] |
| MnS [30] |
| Cd-S-Se [31] |
| ZnSe [32] |
| NiSe [33, 34] |
| PbSe [35] |
| Cd-Zn-Te [36] |
| SnSe [37] |
| FeS [38] |
| CuS [39] |
| Zn-Ni-S [40] |
| Ni-Pb-S [41] |
| In ₂ S ₃ [42] |
| MnS ₂ [43] |
| Ag-Sn-S [44] |
| FeS [45] |
| CdSe[46] |
| PbS [47] |

As pointed out by many researchers, there are many disadvantages could be observed in dye sensitized solar cell including higher recombination of charge carriers between the nanoparticle grain boundaries of photoanode, dye agglomeration over photoanode and the liquid electrolyte degradation due to environmental factors [19]. Therefore, another type of solar cell called thin film solar cell could be used to convert sunlight into electrical energy. Recently, thin films solar cell is produced by depositing thin layers on the substrate using chemical deposition technique. Many researchers have reported that the chemical bath deposited films (Table 1) and electrodeposited thin films (Table 2) always been cheaper but less

efficient than conventional crystalline silicon technology.

TABLE II. ELECTRO DEPOSITED THIN FILMS

| |
|---|
| CdSe [48] |
| ZnS [49] |
| CuInSe ₂ [50] |
| Cu ₄ SnS ₄ [51] |
| Cu ₂ ZnSnS ₄ [52] |
| Cu _x Sn _y S _z [53] |
| ZnTe [54] |

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CONCLUSION

Literature review shows that there are many scientists actively engaged in the research of dye sensitized solar cell. Dye sensitized solar cells could be fabricated based on the TiO₂, CuO, NiO and ZnO acting as an electrode. This type of solar cell has potential to be renewable energy sources due to can convert the solar energy into electrical energy. This review paper clearly shows that the conversion efficiency depends on the photo electrodes.

REFERENCES

- [1] T. Pauporte, and C. Magne, "Impedance spectroscopy study of N719 sensitized ZnO based solar cells," *Thin Solid Films*, vol. 560, pp. 20-26, 2014.
- [2] P. Srivastava, and L. Bahadur, "Dye sensitized solar cell based on nanocrystalline ZnO thin film electrodes combined with a novel light absorbing dye Coomassie Brilliant Blue in acetonitrile solution," *Int. J. Hydrogen Energy*, vol. 37, pp. 4863-4870, 2012.
- [3] W.H. Thomas, A.J. Rebecca, B.F.M. Alex, V.R. Hal, and T.H. Joseph, "Advancing beyond current generation dye sensitized solar cells," *Energy Environ. Sci.*, vol. 1, pp. 66-78, 2008.
- [4] M. Hamadian, J. Ghomi, M. Hosseinpour, R. Masoomi, and V. Jabbari, "Uses of new natural dye photosensitizers in fabrication of high potential dye sensitized solar cells (DSSCs)," *Mater. Sci. Semicond. Processing*, vol. 27, pp. 733-739, 2014.
- [5] L. Yu, J. Yang, and Y. Meng, "Nanostructured ZnO thin films by SDS-assisted electrodeposition for dye sensitized solar cell applications," *Ceram. Int.*, vol. 39, pp. 5049-5052, 2013.
- [6] U. Mehmood, S. Rahman, K. Harrabi, I.A. Hussein, and B.V.S. Reddy, "Recent advances in dye sensitized solar cells," *Adv. Mater. Sci. Technol.*, <http://dx.doi.org/10.1155/2014/974782>, 2014.
- [7] T. Ganesh, S.B. Sambhaji, S.M. Rajaram, and S. Han, "Crystallographic phase mediated dye sensitized solar cell performance of ZnO nanostructures," *Scr. Mater.*, vol. 69, pp. 291-294, 2013.
- [8] K. Suman, and B. Lal, "Characterization of some metal free organic dyes as photosensitizer for nanocrystalline ZnO based dye sensitized solar cells," *Int. J. Hydrogen Energy*, vol. 36, pp. 11620-11627, 2011.
- [9] A.A. Rani, and S. Ernest, "Structural, morphological, optical and compositional characterization of spray deposited Ga doped ZnO thin film for dye sensitized solar cell application," *Superlattices Microstruct.*, vol. 75, pp. 398-408, 2014.
- [10] C. Chou, F. Chou, Y. Ding, and P. Wu, "The effect of ZnO-coating on the performance of a dye sensitized solar cell," *Sol. Energy*, vol. 86, pp. 1435-1442, 2012.
- [11] D. Anca, P. Gabriel, and M. Florin, "Investigations on the influence of surfactant in morphology and optical properties of zinc oxide nanopowders for dye-sensitized solar cells applications," *Mater. Sci. Semicond. Processing*, vol. 16, pp. 1095-1104, 2013.
- [12] Q. Wu, Y. Shen, L. Li, M. Cao, F. Gu, and L. Wang, "Morphology and properties of NiO electrodes for p-DSSCs based on hydrothermal method," *Appl. Surf. Sci.*, vol. 276, pp. 411-416, 2013.
- [13] P. Suri, M. Panwar, and R.M. Mehra, "Photovoltaic performance of dye sensitized ZnO solar cell based on Eosin-Y photosensitizer," *Mater. Sci.-Poland*, vol. 25, pp. 137-144, 2007.
- [14] S. Suphadate, and M. Sasimonton, "Additive SnO₂-ZnO composite photoanode for improvement of power conversion efficiency in dye sensitized solar cell," *Proc. Manuf.*, vol. 2, pp. 108-112, 2015.
- [15] K.S. Jitendra, M.S. Akhtar, S. Ameen, P. Srivastava, and G. Singh, "Green synthesis of CuO nanoparticles with leaf extract of *Calotropis gigantea* and its dye-sensitized solar cells applications," *J. Alloys Compd.*, vol. 632, pp. 321-325, 2015.
- [16] H. Masood, G. Afshar, and J. Vahid, "High performance dye sensitized solar cells (DSSCs) achieved via electrophoretic technique by optimizing of photoelectrode properties," *Mater. Sci. Semicond. Processing*, vol. 16, pp. 1352-1359, 2013.
- [17] S.A. Mozaffari, M. Ranjbar, E. Kouhestanian, H.S. Amoli, and M.H. Armanmehr, "An investigation on the effect of electrodeposited nanostructured ZnO on the electron transfer process efficiency of TiO₂ based DSSC," *Mater. Sci. Semicond. Processing*, vol. 40, pp. 285-292, 2015.
- [18] C. Chou, Y. Lin, R. Yang, and K. Liu, "Preparation of TiO₂/NiO composite particles and their

applications in dye-sensitized solar cells,” *Adv. Powder Technol.*, vol. 22, pp. 31-42, 2011.

[19] R. Parthiban, D. Balamurugan, and B.G. Jeyaprakash, “Spray deposited ZnO and Ga doped ZnO based DSSC with bromophenol blue dye as sensitizer: Efficiency analysis through DFT approach,” *Mater. Sci. Semicond. Processing*, vol. 31, pp. 471-477, 2015.

[20] K. Anuar, S.M. Ho, W.T. Tan, and C.F. Ngai, “Influence of triethanolamine on the chemical bath deposited NiS thin films,” *Am. J. Appl. Sci.*, vol. 8, pp. 359-361, 2011.

[21] H. Taz, R. Ruther, A. Malasi, S. Yadavali, C. Carr, J. Nanda, and R. Kalyanaraman, “In situ localized surface plasmon resonance (LSPR) spectroscopy to investigate kinetics of chemical bath deposition of CdS thin films,” *J. Phys. Chem. C.*, vol. 119, pp. 5033-5039, 2015.

[22] K. Anuar, N. Saravanan, W.T. Tan, S. Atan, Z. Kuang, M.J. Haron, and S.M. Ho, “Effect of Deposition Period and pH on Chemical Bath Deposited Cu_4SnS_4 Thin Films,” *Phil. J. Sci.*, vol. 138, pp. 161-168, 2009.

[23] L. Zhou, Y. Li, and Y. Dong, “Preparation and characterization of $\text{Cd}_{1-x}\text{Zn}_x\text{S}$ thin films with chemical bath deposition,” *Adv. Mater. Res.*, vol. 1088, pp. 86-90, 2015.

[24] K. Anuar, S.M. Ho, K.S. Lim, and N. Saravanan, “SEM, EDAX and UV-Visible studies on the properties of Cu_2S thin films,” *Chalcogenide Lett.*, vol. 8, pp. 405-410, 2011.

[25] V.N. Kumar, R. Suryakarthick, S. Karuppusamy, M. Gupta, Y. Hayakawa, and R. Gopalakrishnan, “Effect of precursor concentration on the properties and tuning of conductivity between p-type and n-type $\text{Cu}_{1-x}\text{Cd}_x\text{S}_2$ thin films deposited by a single step solution process as a novel material for photovoltaic applications,” *RSC Adv.*, vol. 5, pp. 23015-23021, 2015.

[26] K. Anuar, W.T. Tan, M. Jelas, S.M. Ho, and S.Y. Gwee, “Effects of deposition period on the properties of FeS_2 thin films by chemical bath deposition method,” *Thammasat Int. J. Sci. Technol.*, vol. 15, pp. 62-69, 2010.

[27] M. Safonova, E. Mellikov, V. Mikli, N. Revathi, and N. Volobujeva, “Chemical bath deposition of SnS thin films from the solutions with different concentrations of tin and sulphur,” *Adv. Mater. Res.*, vol. 1117, pp. 183-186, 2015.

[28] K. Anuar, S.M. Ho, S. Atan, and M.J. Haron, “The effect of the pH value on the growth and properties of chemical bath deposited SnS thin films,” *Res. J. Chem. Environ.*, vol. 15, pp. 45-48, 2011.

[29] M.G. Claudia, P.A. Luque, A. Beltran, A.R. Nestor, E. Lugo, A. Castillo, M.A. Lopez, and A. Olivas, “Study of morphology of ZnS thin films

deposited on different substrates via chemical bath deposition,” *Scanning*, doi: 10.1002/sca.21227, 2015.

[30] K. Anuar, and S.M. Ho, “Deposition and characterization of MnS thin films by chemical bath deposition method,” *Int. J. Chem. Res.*, vol. 1, pp. 1-5, 2010.

[31] E.A. Ramirez, M.A. Perez, J.R. Hernandez, and G.C. Puente, “Bath atomic composition and deposition time influence on the properties of nanostructured $\text{CdS}_{0.5}\text{Se}_{0.5}$ thin films synthesized by CBD,” *Mater. Chem. Phys.*, vol. 165, pp. 119-124, 2015.

[32] K. Anuar, S.M. Ho, W.T. Tan, Kelvin, and N. Saravanan, “Composition, morphology and optical characterization of chemical bath deposited ZnSe thin films,” *Eur. J. Appl. Sci.*, vol. 3, pp. 75-80, 2011.

[33] G. Nader, and A.B. Tahere, “Shapes alteration and optical band gap controlling in NiSe nanostructure thin films with deposition temperature changing,” *Optik-Int. J. Light Electron Opt.*, vol. 126, pp. 4557-4560, 2015.

[34] K. Anuar, W.T. Tan, A.H. Abdullah, H.M. Jelas, N. Saravanan, S.M. Ho, and M. Yazid, “Chemical bath deposition of NiSe thin films from alkaline solutions using triethanolamine as complexing agent,” *Oriental J. Chem.*, vol. 25, pp. 813-816, 2009.

[35] K. Anuar, S.M. Ho, and N. Saravanan, “Preparation of lead selenide thin films by chemical bath deposition method in the presence of complexing agent (tartaric acid),” *Turk. J. Sci. Technol.*, vol. 6, pp. 17-23, 2011.

[36] S.N. Vidhya, O.N. Balasundaram, and M. Chandramohan, “The effect of annealing temperature on structural, morphological and optical properties of CdZnTe thin films,” *Optik-Int. J. Light Electron Opt.*, vol. 126, pp. 5460-5463, 2015.

[37] K. Anuar, S.M. Ho, W.T. Tan, S.M. Ho and N. Saravanan, “Temperature-dependent surface topography analysis of SnSe thin films using atomic force microscopy,” *Asian J. Res. Chem.*, vol. 5, pp. 291-294, 2012.

[38] S.A. Muhammad, A. Asma, and A.M. Mohammad, “Synthesis of mackinawite FeS thin films from acidic chemical baths,” *Mater. Sci. Semicond. Processing*, vol. 32, pp. 1-5, 2015.

[39] K. Anuar, S.M. Ho, W.T. Tee, K.S. Lim, and N. Saravanan, “Morphological characterization of CuS thin films by atomic force microscopy,” *Res. J. Appl. Sci. Eng. Technol.*, vol. 3, pp. 513-518, 2011.

[40] S. Reza, and D. Soraya, “An investigation on optical characteristics of nanocrystalline ZnS:Ni thin films prepared by chemical deposition method,” *Spectrochim. Acta Part A: Molecular and Biomolecular Spectroscopy*, vol. 149, pp. 941-948, 2015.

[41] S.M. Ho, “Influence of complexing agent on the growth of chemically deposited $\text{Ni}_3\text{Pb}_2\text{S}_2$ thin

films," *Oriental J. Chem.*, vol. 30, pp. 1009-1012, 2014.

[42] F. Mesa, W. Chamorro, and M. Hurtado, "Optical and structural study of In_2S_3 thin films growth by co-evaporation and chemical bath deposition on Cu_3BiS_3 ," *Appl. Surf. Sci.*, vol. 350, pp. 38-42, 2015.

[43] K. Anuar, A.H. Abdullah, S.M. Ho, and N. Saravanan, "Influence of deposition time on the properties of chemical bath deposited manganese sulfide thin films," *Av. Quim.*, vol. 5, pp. 141-145, 2010.

[44] L. Yeh, and K. Cheng, "Preparation of chemical bath synthesized ternary Ag-Sn-S thin films as the photoelectrodes in photoelectrochemical cell," *J. Power Sources*, vol. 275, pp. 750-759, 2015.

[45] K. Anuar, S.M. Ho, S. Atan, and N. Saravanan, "X-ray diffraction and atomic force microscopy studies of chemical bath deposited FeS thin films," *Studia UBB. Chemia*, vol. 55, pp. 5-11, 2010.

[46] A.V. Cephas, S.D. Sartale, J.M. Patil, K.P. Ghoderao, J.P. Sawant, and R.B. Kale, "Room temperature chemical bath deposition of cadmium selenide, cadmium sulfide and cadmium sulfoselenide thin films with novel nanostructures," *Solid State Sci.*, vol. 48, pp. 186-192, 2015.

[47] Y. Ersin, Y. Yasin, and B. Buse, "Optimization of synthesis conditions of PbS thin films grown by chemical bath deposition using response surface methodology," *J. Alloys Compd.*, vol. 642, pp. 63-69, 2015.

[48] S. Mahato, and A.K. Kar, "Structural, optical and electrical properties of electrodeposited cadmium selenide thin films form applications in photo detector and photoelectrochemical cell," *J. Electroanal. Chem.* vol. 742, pp. 23-29, 2015.

[49] K. Anuar, N. Saravanan, S.M. Ho, and K. Noraini, "XRD and AFM studies of ZnS thin films produced by electrodeposition method," *Arabian J. Chem.*, vol. 3, pp. 243-249, 2010.

[50] D. Pottier, and G. Maurin, "Preparation of polycrystalline thin films of CuInSe_2 by electrodeposition," *J. Appl. Electrochem.*, vol. 19, pp. 361-367, 1989.

[51] K. Anuar, K. Zulkefly, S. Atan, H. Jelas, W.T. Tan, and S.M. Ho, "Effects of deposition potential on Cu_4SnS_4 thin films prepared by electrodeposition technique," *Arabian J. Sci. Eng.*, vol. 35, pp. 83-92, 2010.

[52] M. Farinella, R. Inguanta, T. Spano, P.M. Livreri, S. Piazza, and C. Sunseri, "Electrochemical deposition of CZTS thin films on flexible substrate," *Energy Proc.*, vol. 44, pp. 105-110, 2014.

[53] D.B. Francesco, B. Ilaria, C. Stefano, C. Serena, D.L. Antonio, L. Alessandro, V. Francesco, M.M. Maurizio, L.F. Maria, and I. Massimo, "Electrodeposition of ternary $\text{Cu}_x\text{Sn}_y\text{S}_z$ thin films for photovoltaic applications," *Prog. Photovoltaics*, vol. 22, pp. 97-106, 2014.

[54] O. Skhouni, A.E. Manouni, M. Mollar, R. Schrebler, and B. Mari, "ZnTe thin films grown by electrodeposition technique on fluorine tin oxide substrates," *Thin Solid Films*, vol. 564, pp. 195-200, 2014.