



## COPPER (I) OXIDE (Cu<sub>2</sub>O) BASED SOLAR CELLS - A REVIEW

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### ABSTRACT

**Copper (I) oxide (Cu<sub>2</sub>O) is a potential material for the fabrication of low cost solar cells for terrestrial application. A detailed survey on the previous work so far carried out on Cu<sub>2</sub>O based solar cells has been presented. The aspects discussed include the fabrication of Schottky (metal/semiconductor) barrier Cu<sub>2</sub>O solar cells, where different low work function metals are used to form the Schottky barrier solar cells. The problems associated with the Cu<sub>2</sub>O Schottky barrier solar cells and efforts made at improving the performance of these solar cells are highlighted. Discussions on heterojunction solar cells with Cu<sub>2</sub>O have also been presented. Various transparent conducting oxides (TCOs) used in forming the heterojunction cells are mentioned, and successes and failures of using them highlighted. The technological developments of these cells are still in their infancy and the performance remains very poor. The root causes of this poor performance are analyzed and possible areas for future research in the field are outlined.**

**Keyword: Copper (I) oxide, Solar cells, Solar cells, Schottky barrier solar cells**

### INTRODUCTION

The world's major energy sources are non-renewable, and are faced with ever increasing demand, thus are not expected to last long. Besides being non-renewable, these sources, mainly of fossil fuels, contribute tremendously to the perennial problem of global warming. The eminent depletion and pollution problems of the above energy sources make the international community focus attention on alternative sources of energy, and solar energy appears highly promising. This energy is emitted from the sun primarily as electromagnetic radiation in the ultraviolet to infrared and radio spectral regions (0.2 to 3 $\mu$ m). The sun has a reasonable stable life time with a projected constant radiative energy output of over 10 billions (10<sup>10</sup>) years (Sze, 1981).

A solar cell performs two functions: photogeneration of charge carriers in a light absorbing material and separation of the charge carriers to a conductive contact that will transmit the electricity. In its simplest form, the solar cell consists of a junction formed between n-type and p-type semiconductors, either of the same material (homojunction) or different materials (Schottky or heterojunction).

Solar cells are electronic devices used for the direct conversion of solar energy to electricity, using the photovoltaic (PV) effect. In addition to other advantages, the method of converting solar energy to electricity is pollution free, and appears a good practical solution to the global energy problems; especially if practical economic means of direct conversion can be developed.

Cu<sub>2</sub>O is a non-stoichiometric defect p-type (Walter, 1951) semiconductor and its potential for the design of solar cells have been recognized since 1920. This was before silicon, germanium and other potential semiconducting materials were discovered. The development of these materials for the fabrication of solar cells has reached advanced stages

while that on Cu<sub>2</sub>O, as at the time of writing this paper, has been slow. However, interest in Cu<sub>2</sub>O was revived during the mid seventies by the photovoltaic community as a possible low cost material for solar cells. Several groups (Grondhal, 1933, Lange, 1938, Trivich *et al.*, 1976) became interested in Cu<sub>2</sub>O solar cells in particular. This interest lasted for about eight years during which hardly a dozen research papers were published.

These drawbacks notwithstanding, Cu<sub>2</sub>O remains an attractive alternative to silicon and other semiconductors being favored at present for terrestrial applications. Its advantages are:

- i) it is non-toxic,
- ii) there is abundance of the starting material (which is copper), and
- iii) its production process is simple.

Cu<sub>2</sub>O has a direct band-gap of 2 eV which is suitable for photovoltaic conversion. Theoretical calculations have predicted an electrical power conversion efficiency of approximately 12%. The practical electrical power conversion efficiencies obtained by researchers, in the past, are below 2% (Noguet *et al.*, 1977). However, recently efficiency of 2% has been reported (Mittiga *et al.*, 2006).

The researches carried out during the mid-seventies and early eighties have now helped in revealing some of the mysteries surrounding this material and a perfect understanding of the various causes for the poor performance of Cu<sub>2</sub>O solar cells are now known. The purpose of this review is to survey the work done on Cu<sub>2</sub>O solar cells starting from the seventies to date. In what follows the methods of the production of Cu<sub>2</sub>O shall be discussed. Fabrication of and discussion on Cu<sub>2</sub>O solar cells are presented. Subsequently, a discussion on the performance of these cells is reported. Finally, various conclusions are drawn and suggestions for further work on Cu<sub>2</sub>O solar cells are provided.

### Defects in Cu<sub>2</sub>O

When a crystal of compound XY is formed it is usually thought of as having equal numbers of X and Y atoms. Such a crystal is said to be stoichiometric. However, this stoichiometry is not obeyed by many solids as the ratio of atoms in them is slightly different from the ratio in one mole. Such non-stoichiometric compounds balance their structures by the presence of defects: vacancies, interstitials or both (Sear and Fortin, 1984). Cuprous oxide is one of such non-stoichiometric materials with formula Cu<sub>2-δ</sub>O. The deviation from stoichiometry, δ is generally attributed to some imperfections. Sears *et al.* 1984 reported that an excess of oxygen, as a result of stoichiometry, is the major active impurity and gives a p-doped semiconductor.

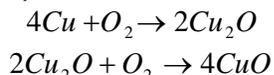
### Methods of Production Of Cu<sub>2</sub>O

The most important methods for the production of Cu<sub>2</sub>O are by thermal oxidation, electrodeposition and by sputtering.

#### Thermal Oxidation

This is by far the most widely used method of producing Cu<sub>2</sub>O for the fabrication of solar cells. The procedure involves the oxidation of high purity copper at an elevated temperature (1000 – 1,500 °C) for times ranging from few hours to few minutes depending on the thickness of the starting material (for total oxidation) and the desired thickness of Cu<sub>2</sub>O (for partial oxidation).

The oxidation process can be carried out either in pure oxygen or in laboratory air. Cu<sub>2</sub>O has been identified to be stable at limited ranges of temperatures and oxygen pressure. It has been indicated that during oxidation, Cu<sub>2</sub>O is formed first and after a sufficiently long oxidation time, CuO is formed. However, at temperatures below 1000°C and at atmospheric pressure, mixed oxides of Cu<sub>2</sub>O and CuO are formed as observed from the X-ray diffraction (XRD) results. It has been suggested that the probable reactions that could account for the presence of CuO in layers oxidized below 1000°C are (Musa *et al.*, 1998):



The unwanted CuO can be removed using an etching solution containing FeCl<sub>3</sub>, HCl and NaCl .

The oxidation process is followed by annealing the sample at 500 °C and then stopping the process by quenching in cold water. This process leads to good quality polycrystalline Cu<sub>2</sub>O with the bulk resistivity in the range of 10<sup>2</sup> -10<sup>4</sup> ohmcm (Trivich *et al.*, 1978 and Economou *et al.*, 1982). The resistivity can further be lowered by oxidizing in the presence of chlorine gas. Resistivities below 100 ohmcm have been reported in the literature (Olsen *et al.*, 1980 and Olsen *et al.*, 1982) using this procedure.

It is also worthy of note that the purity of the starting Cu<sub>2</sub>O material can have a significant impact on the quality of Cu<sub>2</sub>O and the performance of the resulting solar cell. A number of pre- and post-oxidation treatments have been suggested in the literature (Olsen *et al.*, 1982) which involves cleaning,

etching, polishing and annealing the material prior to and after the oxidations.

### Electro-deposition

Another method of producing thin films of Cu<sub>2</sub>O is by electro-deposition. Thin films of Cu<sub>2</sub>O can be electro-deposited by cathodic reduction of an alkaline cupric lactate solution, either on metallic substrates or on transparent conducting glass slides coated with highly conducting semiconductors (Noguét *et al.*, 1977).

The properties of the electrodeposited films of Cu<sub>2</sub>O are largely similar to those prepared by thermal oxidation. The grain sizes of the electrodeposits varies from 0.1 to 10µm. The major problem, however, is in the high resistivity (10<sup>4</sup> – 10<sup>6</sup> ohmcm) of the electrodeposited Cu<sub>2</sub>O film ( Economou *et al.*, 1982).

### Sputtering

Cathode sputtering is essentially one of the methods used for the preparation of thin films. The method requires very low pressure in the working space and therefore makes use of vacuum technique. The material to be sputtered is used as a cathode in the system in which a glow discharge is established in an inert gas at a pressure of 10<sup>-1</sup> – 10<sup>-2</sup> torr and a voltage of a few kilovolts. The substance on which the film is to be deposited is placed on the anode of the system.

The positive ions of the gas created by the discharge are accelerated towards the cathode (target). Under the bombardment of the ions the material is removed from the cathode (mostly in the form of neutral atoms or in the form of ions). The liberated components condense on surrounding areas and consequently on the substrates placed on the anode.

Reactive sputtering is used in the production of Cu<sub>2</sub>O. A chemical reaction that occurs with the cathode material (Cu in this case) by the active gas (oxygen) either added to the working gas or as the working gas itself. The resistivity of the deposited Cu<sub>2</sub>O film can be controlled over a wide range by simply varying the oxygen pressure. Cu<sub>2</sub>O films of resistivity as low as 25 ohmcm have been reproducibly obtained (Drobny *et al.*, 1979) by this technique.

### Fabrication of Cu<sub>2</sub>O Solar Cells

Even though Cu<sub>2</sub>O was known since 1904, very little attention was paid to it. The few research works done on it were mainly concerning its photoresponse, as reported (Olsen *et al.*, 1982 and Wikipedia, 2008). As a result of pioneering work in 1975 at National Science foundation and at the Joint Centre for Graduate Studies (JCGS) on fabrication and characterization of Cu<sub>2</sub>O cells that Cu<sub>2</sub>O solar cells of 1% efficiency were fabricated in 1978. Further works on the cell at JCGS yielded an efficiency of 1.8%. Experimental and theoretical studies conducted at JCGS group highlighted more on the potential viability of Cu<sub>2</sub>O for low cost photovoltaic (PV) power generation.

Until recently, efforts in the past to fabricate efficient solar cells were confined to Schottky barrier solar cells (Herion et al, 1980 and Olsen et al, 1982). However, in recent times, single polycrystalline Cu<sub>2</sub>O used for solar cells are obtained via any of the methods presented earlier in this paper. The Schottky barrier solar cells and the heterojunctions have been extensively investigated. Also, in the past, workers have not succeeded in producing n – type Cu<sub>2</sub>O and therefore a homojunction cell structure (Trivich, 1982) could not be fabricated. Recently, Cu<sub>2</sub>O homojunction solar cells have been fabricated (Longcheng and Meng, 2007 and Kunhee and Meng, 2009).

#### **Schottky Barrier Cu<sub>2</sub>O Solar Cells**

A frontwall Schottky barrier solar cell (Olsen et al., 1980) is usually fabricated by evaporating a metal on top of Cu<sub>2</sub>O in high vacuum chambers. A number of metals on Cu<sub>2</sub>O have been tried and the best performance has been obtained with a natural Cu/Cu<sub>2</sub>O junction.

A new method of obtaining Cu<sub>2</sub>O has been reported (Iwanoski et al., 1985). This technique uses hydrogen ion bombardment of Cu<sub>2</sub>O surface thus reducing the top surface to copper. Schottky barrier solar cells can also be fabricated in the back wall structure. This mode requires a natural junction using copper or any other material as the base and depositing a layer of Cu<sub>2</sub>O on top of the metal base. The light then illuminates the junction through the semiconductor side. But back wall cells require thin layer of Cu<sub>2</sub>O ( $\leq 1\mu\text{m}$ ) because of the high absorption coefficient and low diffusion length of the minority carriers (Trivich et al., 1978). Partial thermal oxidation can be used to produce the thin layer. Back wall solar cells of Cu<sub>2</sub>O offer greater advantage over front wall cells because of the mechanical stability of the base materials and simplicity of fabrication.

#### **Heterojunction Solar Cells**

A heterojunction solar cell is fabricated by depositing n-type semiconductor of suitable band gap on Cu<sub>2</sub>O. Methods like vacuum deposition, sputtering and electrodeposition have been used for the deposition. Several heterojunction solar cell structures have been studied and reported (Herion et al., 1980). Examples are the zinc oxide- cuprous oxide (ZnO/Cu<sub>2</sub>O) and cadmium oxide-cuprous oxide (CdO/Cu<sub>2</sub>O) solar cells.

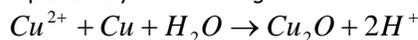
Transparent conducting metal oxides, being n-type were used extensively in the production of heterojunction cells using p-type Cu<sub>2</sub>O. Herion et al. (1980) have reported on a fairly detailed study of ZnO/Cu<sub>2</sub>O devices. This was achieved due to the interest they had on metal oxides being generally stable compounds and the assumption that they are not likely to react with Cu<sub>2</sub>O. Eventually, the cell exhibited poor performance. The cell characteristics were clearly influenced by the copper rich region adjacent to Cu<sub>2</sub>O substrate. It was finally concluded that ZnO/Cu<sub>2</sub>O heterojunction is essentially Cu/Cu<sub>2</sub>O Schottky cell since Zn reduces Cu<sub>2</sub>O to Cu. Hence reaction occurs with this type of cells too. Indium Tin Oxide (ITO) devices prepared by different techniques on Cu<sub>2</sub>O were also reported (Herion et al., 1980 and Georgieva and Ristov, 2002). In all cases resistivity

was found to be high and efficiency very low. Tanaka et al. (2004) have reported an intensive work on Cu<sub>2</sub>O prepared with TCOs. They used TCO thin films such as indium oxide (In<sub>2</sub>O<sub>3</sub>), tin oxide (SnO<sub>2</sub>) and multicomponent oxides like aluminium- zinc-oxide(AZO) and aluminium- zinc- indium- tin- oxide (AZITO), in addition to ZnO. Highest efficiency of 1.2 % was obtained with AZO-Cu<sub>2</sub>O devices prepared at 150 °C and measured under air mass 2 (AM2) illumination. Others yielded disappointing results. A better cell was reported by Trivich et al. (1981) using CdO. It was found to generate an open circuit voltage,  $V_{oc} = 0.4\text{V}$  and a short circuit current,  $I_{sc} = 2\text{mA cm}^{-2}$ .

There was no reduction of the Cu<sub>2</sub>O layer observed, which shows the absence of any chemical reaction at the CdO/Cu<sub>2</sub>O interface. This suggested a possibility of avoiding chemical reaction at the interface by the use of heterojunction, especially, of oxides on Cu<sub>2</sub>O. However, the best solar cell, to date using TCO thin films is a multicomponent oxides, ITO/ZnO/Cu<sub>2</sub>O solar cell. It was reported to have an efficiency of 2% (Mittiga et al., 2006). Copper (1) sulphide (Cu<sub>2</sub>S) is another promising material for heterojunction devices for the formation of n-Cu<sub>2</sub>O/Cu<sub>2</sub>S heterojunction solar cell since Cu<sub>2</sub>S is a p-type semiconductor as grown. Works in the past were aimed at fabricating CdS/Cu<sub>2</sub>S heterojunction devices. Fajinmi, 2000, reported on the deposition of Cu<sub>2</sub>S, while Varkey, 1990, reported on copper (1) sulphide/crystalline silicon (Cu<sub>2</sub>S/c-Si) solar cells.

#### **Homojunction Solar Cells**

In the past the low efficiency of Cu<sub>2</sub>O cells was attributed to the lack of n-type Cu<sub>2</sub>O, since an approach to achieving n-type doping has not yet been fully developed. Without it, the early studies had to rely on Schottky junctions and p-n heterojunctions for photovoltaic devices, which do not provide high efficiency. However, Fernando et al, 2002, has reported the possibility of obtaining n-type photo-responses from clean copper plates, immersed in CuSO<sub>4</sub> solution for a few days. Subsequently, the same researchers reported the n-type Cu<sub>2</sub>O produced by heating copper sheets in CuSO<sub>4</sub> solution. The formation of the n-type Cu<sub>2</sub>O on copper surface, by heating copper sheets in CuSO<sub>4</sub> solution, can be explained by the following chemical reaction:



The long held consensus is that the best approach to improve cell efficiency in Cu<sub>2</sub>O-based photovoltaic devices is to achieve both p- and n-type Cu<sub>2</sub>O and thus p-n homojunction of Cu<sub>2</sub>O solar cells. Experimental results in achieving both conduction types in electrochemically deposited Cu<sub>2</sub>O by varying solution pH was reported by Longcheng and Meng, 2007. This enables p-type and n-type Cu<sub>2</sub>O to be deposited electrochemically in sequence to form a p-n homojunction of Cu<sub>2</sub>O. However, the first homojunction solar cell of Cu<sub>2</sub>O using the electrochemical method was made by Kunhee and Meng (2009). The cell has only 0.1% efficiency due to high resistivity of the p- and n-Cu<sub>2</sub>O layers (or substrate).

### Performance of Cu<sub>2</sub>O Solar Cells

The observed conversion efficiency of Cu<sub>2</sub>O cells has remained far below the theoretical value, regardless of the method of growth of Cu<sub>2</sub>O and the mode of fabrication of the cells. The best results obtained so far are in the range of 1-2 %. Many reasons have been advanced for this low performance.

Barrier height measurements in various Schottky barrier solar cells have shown that values are always in the range 0.7-0.9 eV regardless of the metal except for the case of gold and silver which form ohmic contacts with Cu<sub>2</sub>O. This apparent plateau for the value of barrier heights is believed to be the principal cause of the low performance of the Cu<sub>2</sub>O Schottky barrier solar cells. Studies on Schottky barrier solar cell indicate that there always exist a copper rich region at the interface between metal and Cu<sub>2</sub>O regardless of the choice of the metal used. All Schottky type cells are therefore essentially Cu/Cu<sub>2</sub>O solar cells and hence the constancy of the value of barrier height and the low electrical power conversion efficiency.

Papadimitriou *et al.* (1981) have reported the results of their study on ZnO/Cu<sub>2</sub>O junction solar cells. The best values they obtained for open – circuit voltage ( $V_{oc}$ ) were of the order of 0.3 V. A copper rich region adjacent to the Cu<sub>2</sub>O substrate was found to be responsible for dictating the cell characteristics. For the case of CdO/Cu<sub>2</sub>O heterojunction formed at room temperature, no copper metal was found at the interface. The CdO/Cu<sub>2</sub>O cell showed a short circuit current of 2 mAcm<sup>-2</sup> and  $V_{oc}$  of 0.4 V.

The method of electrodeposition is particularly attractive for its simplicity, low cost and possibility of making large area thin films (Wijesundara *et al.*, 2006). But resistivity of electrodeposited Cu<sub>2</sub>O films were reported to be quite high, of the order of 10<sup>4</sup> – 10<sup>6</sup> Ωcm (Economou *et al.* 1982 and Wijesundara *et al.*, 2000), and therefore, the photovoltaic properties of the solar cells made with these films were poor. Several other production techniques of Cu<sub>2</sub>O layer were later employed in order to improve on its resistivity. The various methods include thermal oxidation (Sear and Fortin, 1984, Musa, 1995 and Mittiga *et al.*, 2006), chemical vapour deposition, or radio frequency magnetron sputtering (Akimoto *et al.*, 2006). However, little improvements were recorded as the efficiency was still very low, less than 2%.

Surface analyses combined with barrier height studies (Olsen *et al.*, 1979, Herion *et al.*, 1980 and Olsen *et al.*, 1982) indicate that Cu<sub>2</sub>O Schottky barriers made with low work function metals are essentially Cu/Cu<sub>2</sub>O cells due to reduction of the Cu<sub>2</sub>O surface. The copper rich region essentially determines the barrier height. Auger depth profiles showed the occurrence of chemical reaction of thermodynamically more stable oxides at the interface and correspondingly to some reduction of the Cu<sub>2</sub>O. Thallium is reported to be the only metal identified that would not reduce Cu<sub>2</sub>O (Olsen *et al.*, 1982). With 3.7 eV as its work function and theoretical value of a dark current,  $J_0$  of the order of 10<sup>-8</sup> Acm<sup>-2</sup>, a TI/Cu<sub>2</sub>O Schottky barrier has a very large theoretical efficiency. Efficiency of 10% at air mass one (AM1) would be

possible. However, deposition of TI of reasonable sheet conductance is not possible on a substrate at room temperature. It was reported that TI of adequate sheet conductance was deposited on cool substrate. Studies on the TI/Cu<sub>2</sub>O barrier represented only a slight improvement over the Cu/Cu<sub>2</sub>O device. In fact studies on a number of metal/Cu<sub>2</sub>O and metal/insulator/Cu<sub>2</sub>O contacts (Trivich *et al.*, 1981) showed that the barrier heights did not depend on the work function of metal.

Intensive work was done regarding deposition of Cu<sub>2</sub>O and the dopant impurities (Musa, 1995 and Akimoto *et al.*, 2006) but general observations show that the various dopants used to further lower the resistivity showed no significant improvement, with Cadmium as an exception. Nitrogen acts as a p-type dopant. Other methods of improving on the resistivity of the layer were then employed. Annealing of the Cu<sub>2</sub>O layer at moderate temperatures, as reported by Noguét *et al.* (1977); Musa *et al.* (1998) and Wijesundara *et al.* (2006), showed some improvements. Another treatment followed; which is potassium cyanide (KCN) treatment as reported by Akimoto *et al.*, (2006). The above treatments yielded little improvement since the efficiency, as at the time of this report, is still not more than 2%, for Schottky barrier Cu<sub>2</sub>O solar cells.

The encouraging aspect of the above treatments is that, they revealed ways for further improvements. It was shown that the PV properties of Cu<sub>2</sub>O Schottky cells are significantly affected by the surface treatment and crystallinity of Cu<sub>2</sub>O (Fernando *et al.*, 2002 and Tanaka *et al.*, 2004). In particular, the deposition method and conditions are important when depositing a thin film on Cu<sub>2</sub>O sheets.

### Concluding remarks and suggestions for further work

Some reasons based on the analysis given can now be advanced for the low value of electrical power conversion efficiency of Cu<sub>2</sub>O based solar cells.

- (1) The high resistivity of the starting material. This problem contributes greatly to the performance of electrodeposited Cu<sub>2</sub>O cells in particular. Further work need to be done to lower the resistivity of electrodeposited Cu<sub>2</sub>O by doping.
- (2) Non-existence of a technique of doping Cu<sub>2</sub>O to get low resistivity n- type semiconductor before now so that conversion efficiencies greater than 2% for p-n homojunction Cu<sub>2</sub>O solar cell could be fabricated. It is suggested that the work of Fernando *et al.*, 2002 on production of n-type Cu<sub>2</sub>O, Longcheng and Meng, 2007 and Kunhee and Meng, 2009, on p-n Cu<sub>2</sub>O homojunction be pursued vigorously for the purpose of obtaining low resistivity Cu<sub>2</sub>O homojunction solar cells. A better understanding of the stoichiometry of Cu<sub>2</sub>O is desirable in this respect.
- (3) A copper rich or oxygen deficient surface which makes all Schottky barriers essentially a Cu/Cu<sub>2</sub>O structure. A suitable method of getting rid of the Cu-enriched region from the interface need to be explored.

It is therefore clear that a better understanding and remedy to the problems enumerated above demands

more basic and applied investigation on  $\text{Cu}_2\text{O}$ .

## REFERENCES

- Akimoto, K. Ishizuka, S. Yanagita, M. Nawa, Y. Paul, K.G. Sakurai, T. (2006): Thin Film Deposition of  $\text{Cu}_2\text{O}$  and Application for Solar Cells. *Solar energy*, 80: 715 – 722.
- Drobny, V.F and Pulfrey, D.L. (1979): Properties of Resistively-Sputtered Copper Oxide Thin Films. *Thin solid films*, 61: 89-98.
- Economou, N.A. Toth, R.S. Komp R.J. and Trivich, D. (1982): Photovoltaic cells of electrodeposited cuprous oxide. 14<sup>th</sup> IEEE Photovoltaic Spec. Conf. Proc. New York: 1180-1185.
- Fajinmi, G.R. (2000): Electrical Characterization of Chemically Deposited  $\text{Cu}_x\text{S}$  Films for Photovoltaic Application. *Nigerian Journal of Physics*, 12 : 25 – 28.
- Fernando, C.A.N, de Silver, P.H.C. Wethasinha, S.K, Dharmadasa, I.M. Delsol, T. and Simmonds, M.C. (2002): Investigation of n-type  $\text{Cu}_2\text{O}$  Layers Prepared by low Cost Chemical Method for Use in Photovoltaic Thin Film Solar Cells. *Renewable energy*, 26: 521 – 529
- Georgieva, V. and Ristov, M.J. (2002): Electrodeposited Cuprous Oxide on Indium Tin Oxide For Solar Applications. *Solar material and solar cells*. 73: 67 – 73.
- Grondahl, L.O. (1933): *Rev. Mod. Phys.* 5: 141.
- Herion, J. Niekisch, E.A. and Scharl, G. (1980): Investigation of Metal Oxide/Cuprous Oxide Heterojunction Solar Cells. *Solar energy materials* 4: 101 – 112
- <http://en.wikipedia.org/wiki/solar-cell>, 05/03/ 2008.
- Iwanoski, R.J. and Trivich, D. (1984):  $\text{Cu}/\text{Cu}_2\text{O}$  Schottky Barrier Solar Cells Prepared by Multistep Irradiation of a  $\text{Cu}_2\text{O}$  Substrate by  $\text{H}^+$  Ions. *Sol. Cells*, 13: 253 – 264.
- Kunhee, H, and Meng, T. (2009): Electrically Deposited p-n Homojunction Cuprous Oxide. *Solar cells. Solar Energy and Solar Cells*, 93: 153-157
- Lange, B. (1938): *Photoelements and their Applications*, Reinhold publishing Corp; New York,
- Longcheng, W. and Meng, T. (2007): Fabrication and Characterization of p-n Homojunctions in Cuprous Oxide by Electrochemical Deposition. *Electrochemical and Solid State Letters*, 10 (9): H248-H250
- Mittiga, A. Salza, E. Sarto, F. Tucci, M. and Vasanthi, R. (2006): Heterojunction Solar Cell with 2% Efficiency based on a  $\text{Cu}_2\text{O}$  Substrate *Applied physics letters*, 88: 163 502-1 – 163502-2
- Musa, A.O. (1995): Development of Thin Film Copper (I) Oxide for Backwall Schottky Barrier Solar Cells, Ph.D Thesis, University of Ilorin, Nigeria. 108-136.
- Musa, A. O. Akomolafe, T. and Carter, M. J. (1998): Production of  $\text{Cu}_2\text{O}$  Solar Cell Material, by Thermal Oxidation and Study of its Physical and Electrical Properties. *Solar energy materials and solar cells*, Pergamon, 51: 3-4.
- Nogueta, C. Tapiero, M. Schwab, C. Zielinger, J.P. Trivich, D. Komp, R.J. Wang, E.Y. and Wang, K. (1977): Cuprous Oxide as a Photovoltaic Converter. *1<sup>st</sup> European community Photovoltaic conference proc.* P. 1170.
- Olsen, L.C. Addis F.W. and Bohara, R.C. (1980): *14<sup>th</sup> IEEE photovoltaic Specialist Conf. Proc., IEEE*, New York, , p. 462.
- Olsen, L.C. Addis, F.W. and Miller, W. (1982): Experimental and Theoretical Studies of  $\text{Cu}_2\text{O}$  Solar Cells. *Sol. Cells*, 7: 247 – 279.
- Olsen, L.C. and Bohara, R. C. (1979): Explanation for Low Efficiency  $\text{Cu}_2\text{O}$  Schottky Barrier Solar Cells. *Applied physics letters*, 34(1): 47 – 49.
- Papadimitriou, L. Economou, N.A. and Trivich, D. (1981): Heterojunction Solar Cells on Cuprous Oxide. *Solar cells*, 3: 73 - 80.
- Sear, W.M. and Fortin, E.J. (1984): Preparation and Properties of  $\text{Cu}_2\text{O}/\text{Cu}$  Photovoltaic Cells. *Solar Energy materials*, 10: 93 – 103.
- Sze, S.M. (1981): *Physics of semiconductor devices*. John Wiley and Sons, Inc.: 790-835.
- Tanaka, H. Shimakawa, T. Miyata, T. Sato, H. Minami, T. (2004): Electrical and Optical Properties of TCO- $\text{Cu}_2\text{O}$  Heterojunction Devices. *Thin solid films*, 469 – 470: 80 – 85.
- Trivich, D., Wang, E.Y. Komp, R.J. and . Ho, F. (1976): 12<sup>th</sup> IEEE Photovoltaic Specialists Conf. Proc. Baton Rouge: p 875.
- Trivich, D., Wang, E.Y. Komp, R.J. Kakar, A.S. (1978) : *13<sup>th</sup> IEEE Photovoltaic Specialist Conf. Proc.* 174
- Trivich, D. Fish, L. and Iwanowski, R.J. (1982): *16<sup>th</sup> IEEE Photovoltaic Specialist Conf. Proc., IEEE*, New York, p. 1072.
- Trivich, D. (1981): *Solar cells*. 3 73 – 80.
- Varkey, A.J. (1990): Photovoltaic Effect of Chemically Deposited  $\text{Cu}_2\text{S}/\text{C-si}$  Heterojunctions. *Solar and wind Technology*, 7(5): 619 – 621.
- Walter, H.B. (1951): The Copper Oxide Rectifier. *Reviews of Modern Physics*, 23: 203 -212.
- Wijesundara, R.P. Hidaka, M. Koga, K. and Siripala, W. (2006): Growth and Characterization of Potentiostatically Electrodeposited  $\text{Cu}_2\text{O}$  and  $\text{Cu}$  Thin Films. *Thin solid films* 500: 241 – 246.
- Wijesundara, R.P., Perera, L.D.R.D. Jayasuriya, K.D., Siripala W., De Silva K.T.L., Samantilleke A.P., Dharmadasa I.M. (2000): *Solar energy materials and solar cells*, 61: 277-286.