# Role of Photo sensitizer-Reductant for Generation of Electrical Energy in Photo galvanic Cell

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#### **Abstract**

We have constructed p-n heterojunction solar sandwich cells with photo sensitizer in a thin film of reluctant placed between a conducting glass coated with indium oxide and a platinum foil. The current-voltage elations of the cells have been measured in the dark and light under both forward and reverse biases.

Key words: Power Point, Fill Factor, Conversion Effiency, Rose Bengal, Glucose, TEA

#### Introduction

Solar cells convert sunlight directly to electricity with acceptable conversion efficiency. They are virtually free of pollution. There are few reports<sup>1-3</sup> on the construction of solar cells utilizing the semi conducting properties of dyes. The photovoltaic effect in a sandwich cell consisting of phenol Safranine in the solid polycrystalline state has been reported earlier<sup>4</sup>, the system generates a photo voltage of 90 mV within a few seconds. The dyes have characteristic properties of aggregation, which cause low efficiencies due to selfquenching. To increase the efficiency and overcome to this problem, we have constructed a sandwich cell with a thin film of phenosafranin in polyvinyl alcohol coated onto conducting glass (containing In<sub>2</sub>O<sub>3</sub>) and a platinum foil. The preliminary results<sup>5</sup> show that the cell on illumination generates a fairly large photo voltage of 515 mV. This interesting result has led us to make a detailed study of thin-film solar cells consisting of dye such as Rose Bengal and reductants such as TEA and Glucose. The results are reported here. A detailed literature survey reveals that different photo sensitizers and reductants have been used in photo galvanic cell<sup>9</sup>-

#### **Material and Methods**

The dyes such as Rose Bengal supplied by Loba Chemicals were twice re-crystallized from ethanolwater solutions. TEA and Glucose (Loba Chemical) were used without further purification. AR grade polyvinyl alcohol (Loba Chemical) was used without further purification. The solutions were prepared with doubly-distilled water. The conducting glass coated with  $In_2O_3$  was supplied by Nesatron (PPG Industries). The thin films (20-30  $\mu$ m thick) of dyes in polyvinyl alcohol were prepared by spreading the solution of dye of known concentration containing a definite amount of polyvinyl alcohol (~0.8 g/10 ml).

The experimental sandwich cell was constructed with semitransparent conducting glass containing a thin film of dye in. polyvinyl alcohol and a platinum foil by using spring clips. The construction of the sandwich cell and the experimental set-up were described in an earlier paper<sup>31</sup>. A tungsten lamp (220V, 300 W) served as light source and had an intensity of 30mW-cm<sup>-2</sup>. The photo voltage and photocurrent were measured with a Keithley model 642 electrometer and a model 196 multimeter, respectively.

## **Results and Discussion**

On illuminating the conducting glass of the sandwich cell containing the photo sensitizer dye, a photo voltage develops, which attains a maximum value within a few minutes. This photo voltage decays gradually in the dark. The growth and decay of the cell photo voltages using different

dyes of different concentrations in the thin film are shown in Fig.1. The open-circuit photo voltage  $(V_{oc})$  increases with increase in $\mu$  concentration (C) of dye till it reaches a maximum for effective conversion of light in electrical energy and follows a relation that is similar to the Freundlich adsorption isotherm, i.e.  $V_{oc} = KC^{l/m}$ , where K is a constant and m is a number greater than one.

Power Conversion Effiency of Photo galvanic Cell: One of the important characteristics of any electrochemical cell is its power conversion effiency. The i-V characteristics of Rose Bengal-TEA and Rose Bengal-Glucose. Photo galvanic cells have been investigated to estimate the power conversion efficiency of the cell. The possible power output from the cell can be obtained from the rectangle of maximum area which can be drawn under i-V curve. The power point (appoint on the curve where the product of potential and current was maximum) in i-V curves were determined and their fill factors were also calculated. These data are summarized in table 2. The efficiency of the Rose Bengal-TEA photo galvanic cell has been calculated to be 1.248 percent, comparable to that of and Rose Bengal -Glucose photo galvanic cell has been calculated to be 0.744 percent. Power point (A point where the product of photocurrent and photo potential is maximum is determined by using i-V curve. Where the fill factor and Conversion efficiency of the cell is calculated by using following formula.

Fill-factor (n) = 
$$\frac{\text{Vpp x ipp}}{\text{Voc x isc}}$$

Where Vpp = photo potential at power point.

ipp = Current at power point.

Voc = potential in open circuit.

isc = Current in short circuit.

Conversion efficiency = 
$$\frac{\text{Vpp x ipp}}{10.4 \text{ Mw}/} \times 100\%$$

Performance of the photogalvanic cell: All the two systems were studied by applying the desired external load to have the potential and current corresponding to power point. The time t1\2 was determined after removing the source of light. It is the time taken in reaching half the value of power. The performance of cells was studied and comparative values are summarized in table 4.

On the basis of the observed data, the Rose Bengal TEA is most efficient from power point of view (Electrochemical Studies of Photosensitizers)The value of K is 3.16 V, The current-voltage relation of the p-n heterojunction cell in the dark is  $I = I_0[1-(V/V_{bi})]$  [exp(eV/kT)-1], where  $V_{bi}$  is the total electrostatic potential for the two semiconductors and k is Boltzmann's constant.

When the junction is forward biased, i.e. V >> (kT/e), then  $I = I_0[1-(V/V_{bi})] [exp(eV/kT)-1]$ , when it is biased in reverse, i.e.  $V \ll (kT/e)$ , then I = $I_0(V/V_{bi})-I_0$ , where  $I_0$  is the dark current for the system. On the other hand, the photocurrent I<sub>L</sub> develops when these systems are illuminated and the current-voltage relation becomes  $I = I_L + I_0$  [1- $(V/V_{bi})$ ] [exp (eV/kT)-1], When the illuminated cell is biased forward and in reverse, then  $I = I_L + I_0$  $[1-(V/V_{bi})]$  exp (eV/kT) and  $I=I_L+I_0$   $(V/V_{bi})-I_0$ , respectively. Linear and semilog plots of the current-voltage relations of the systems are shown in Figs. 2 and 3 for the dark and with light when the junctions are biased both forward and in reverse. All of the parameters are calculated from these plots.

The ln<sub>2</sub>O<sub>3</sub>-coated glass is an n-type semiconductor with a band gap<sup>32</sup> of 3.6eV, whereas phenosafranin behaves as a p-type semiconductor. The reported band gap<sup>33</sup> for phenosafranin is 2.17 eV. Platinum is used only for ohmic contact. The formation of a p-n heterojunction solar cell is initiated on absorption of visible light by the dye molecules. When visible light is incident on these p-nheterojunctions, an excess of electron-hole pairs is generated and the electrons diffuse to the n-region while the holes diffuse to the p-region. In the absence of an externally applied field, the lightinduced current develops a photovoltage across the heterojunction.The conductivities semiconductor dyes in thin films of reductant increase exponentially with temperature according to the relation<sup>6</sup>  $\sigma(T) = \sigma_0 \exp(-E/2kT)$ , where E is the band gap for the process and  $\sigma_0$  is the

extrapolated conductivity at infinite temperature. The band gaps of dyes in a thin film of reductant have been determined directly from conductivity measurements and plotting lno as a function of 1/T. The results for Rose Bengal- TEA and Rose Bengal-Glucose are 1302.0, and 1080.0 mV, respectively. There is an interesting correlation between the activation energies of the dyes and the photo voltages generated in the film containing dyes. The lower the band gap of the dye, the greater is the photo voltage generated in the solar cell containing dye. Lower activation energy means that less energy is required to generate charge carriers in the semiconductors.

The use of dyes in the film increases the efficiency of photo voltage generation by decreasing selfquenching caused by aggregation. This type of solar cell using a polymer of low resistivity is important since its preparation is simple and the cost is low compared to that of other solar cells.

**Mechanism:** On the basis of these observations, a mechanism is suggested for the generation of photocurrent in the photo galvanic cell as:

#### **Illuminated Chamber**

$$Dye \longrightarrow Dye^*$$
 (6.1)

$$Dye^* + R \longrightarrow Dye (semi or leuco) + R^+ (6.2)$$

#### **Platinum Electrode**

$$Dye - \longrightarrow Dye + e -$$
 (6.3)

## **Dark Chamber**

Dye 
$$+e^-$$
 Dye (semi or leuco) (6.4)

$$Dye^{-} + R^{+} \longrightarrow Dye + R$$
 (6.5)

Where Dye, Dye\*, Dye — , R and R+ are the excited form of dye, semi or leuco dye, reductant and oxidized form of the reductant, respectively.

#### Conclusion

Photo galvanic cells are low cost due to the use of a dye, which are cheap and used in minute quantities reductant like Ascorbic acid, is also not that very expensive. So overall working with a photo galvanic cell has lot of scope for its development.

On the basis of observation in the whole study it is concluded that photo galvanic cells are better option for solar energy conversion and storage. Also this system with better electrical output good performance and storage capacity may be used in near future. According to observed photo galvanic effect in all these two systems, Rose Bengal-TEA system was the most efficient in all the ways.

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#### References

- 1. Meier H., Albrecht W. and Tschirwitz U., Angew., *Chem. Int. Edn.*, **11**, 1051 (**1972**)
- 2. Chamberlain G.A., *Mol. Cryst. Liq. Cryst.*, **93**, 369 (**1983**)
- 3. Panayotatos P., Bird G., Sauers R., Piechowski A. and Husain, S., *Sol. Cells.*, **21**, 301 (**1987**)
- 4. Rohatgi Mukherjee K.K., Roy M. and Bhowmik ,B.B., *Sol. Energy*, **31**, 417 (**1983**)
- 5. Bhowmik B.B. and Jana A.K., *Energy-The International Journa*, **13**, 755 (**1988**)
- 6. Sze S.M., Physics of Semiconductor Devices, 2<sup>nd</sup> edn, Wiley Eastern, New Delhi (**1983**)
- 7. Schumacher L.C., Maniche-Afara S. and Dignam M.J., *J. Electrochem. Soc.*, **133**, 716 (**1986**)
- 8. Meier H., J. Phys. Chem., **69**, 719 (**1965**)
- 9. Punjabi P. B., Vardia J., Madhwani S. and Chaudhary S., *J. Power Sources*, **159**, 747–751 (**2006**)
- 10. Madhwani S., Ameta R., Vardia J., Punjabi P. B. and Sharma V. K., *Energy Sources.*, **29**, 721 729 (**2007**)

- 11. Bohrmann Linde C. and Tausch M. W., *J. Chem. Educ.*, **80**, 1471–1473 (**2003**)
- 12. Monat J. E. and Mc Cusker J. K., *J. Amer. Chem. Soc.*, **122**, 4092 (**2000**)
- 13. Schwarzburg K. and Willig F., *J. Phys. Chem.* **103B**, 5743 (**1999**)
- 14. Tennakone K. and Kumara GRR.A., *J. Photochem. Photobiol.*, **117A**, 137 (**1998**)
- 15. Yadav Sushil, Yadav R. D. and Singh Gautam, *Int. J. Chem. Sci.*, **6(4)**, 1960 (**2008**)
- 16. Meena R.C., Gautam Singh and K.M., Gangotri K. M., Afinidad **59**, 253-256 (**2003**)
- 17. Meena, R.C., and R.S. Sindal., Int. J. Chem. Sci., **2**(3), 321-330 (**2004**).
- 18.Ameta, Suresh C., Sadhana Khamesra, Anil K., Chittoro and K.M. Gangotri.,Int. J. Energy Res., 13, 643, (1989).
- 19.Gongotri, K.M., R.C. Meena and Rajni Meena., J. Photochem and photobiol. A: Chem; **123**, 93, (**1999**)
- 20.Gangotri, K.M. and Chhagan Lal., Int. J. Energy Res., **24**, 365, (**2000**).
- 21.Fisher, A.C., L.M. Peter, E.A., Ponomareve, A.B., Walker, and K.G.U. Wijayantha.,J. Phys. Chem. B. **104(5)**, 949-958 (**2000**)
- 22.Zhang, Zhi-Ying. and Chun-yanliu., J. Photochem. and Photobio. A: Chemistry **130**, 139 (**2000**)

- 23.Gratzel, Michael., J. Photochem.and Photobiol. A: Chemistry **164**, 3-14, (**2004**).
- 24. Coralie Houarner-Rassin, Errol Blart, Pierrick Buvat and Fabriceodobel., J. Photochem. and Photobiol A: Chemistry **186**, 135, **(2007)**.
- 25. Bandara J., U.W. Pradeep and R.G.S.J. Bandara., *Solar cells*, **170**, 273 (**2005**)
- 26. Sindal R. S., Gunsaria R. K., Chandra Mahesh and Meena R. C., *The Arabian Journal for science and Engineering*, **31(2A)**, 177 **(2006)**
- 27. Minna Toivola, Lauripeltokorpi, Janne Halme, Peter Lund, *Solar energy materials and solar Cells*, **91**, 1733 (**2007**)
- 28. Kenisarin mural. and Khamid Mahkamov, Renewable and Sustainable energy reviews, **11**, 1913, **(2007)**
- 29. Sindal R. S., Chandra Mahesh and Meena R. C., J. Ind. Council Chem., **25(2)**, 131-136, **(2008)**.
- 30. Sirvi Sonal, Meena Vijay Kumar and Meena R C, *J. Indian Chem.Soc.* **85**, 825 (**2008**)
- 31. Meena R. C., *J. Indian Chem. Soc.*, **85**, 280 (2008)
- 32. Sindal R. S., Chandra M., Kumar V. and Meena R. C., *Energy Sources*, **31(16)**, **(2009)**
- 33. Kumari Manju, Pachwarya Ram Babu and Meena R.C, *Energy Sources*, **31(11)**, **(2009)**

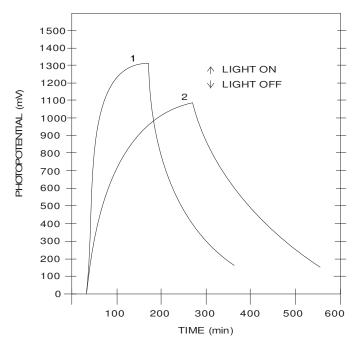


Figure-1: The effect of light on photo voltage generation for solar cells containing dye-reductants mixtures in the thin film. The dyes reductant mixtures

**Table-1: Electrical parameters** 

Observations	Rose Bengal - TEA System	Rose Bengal-Glucose System
Open Circuit voltage (Voc)	1302.0 mV	1080.0 mV
Photopotential (V)	1185.0 mV	968.0 mV
Equilibrium Photocurrent (ieq)	175.0 μΑ	160.0 μΑ
Maximum Photocurrent (imax)	210.0 μΑ	190.0 μΑ
Short circuit current (isc)	175.0 μΑ	160.0 μΑ
Current at power point (ipp)	135.0 μΑ	110.0 μΑ
Potential at power point (Vpp)	962.0 μΑ	704.0 μΑ
Power at power point	68.8 μA/min	62.8 μA/min
Rate of Generation	62.4 µA/min	56.8 μA/min
Conversion Efficiency	1.248%	0.744%
Charging Time	180.0 min	260.0 min
t1/2	90.0 min	60.0 min
Fill factor (n)	0.56	0.44

Table-2: i-V Characteristics of the photo galvanic cells

Systems	Voc (mV)	isc (μA)	Vpp (mV)	ipp (µA)	n
Rose Bengal - TEA System	1302	175.0	962.0	135.0	0.56
Rose Bengal-Glucose System	1080	160.0	704.0	110.0	0.44

The conversion efficiency and sunlight conversion data for these two systems are reported in table 3.

Table-3: Conversion efficiency and sunlight conversion data

Systems	Fill Factor (n)	Conversion Efficiency (%)	Sunlight Conversion Data	
			Photo potential (mV)	Photocurrent ( \mu A)
Rose Bengal-TEA	0.56	1.24	1302	210
Rose Bengal-Glucose	0.44	0.74	1080	190

On the basis of these observations, the highest conversion efficiency was found in Rose Bengal –TEA system.

Table-4: Performance of the photo galvanic cells in dark

System	Power (µW)	t1/2 (min)
Rose Bengal-TEA	68.8	90.0
Rose Bengal-Glucose	62.2	60.0

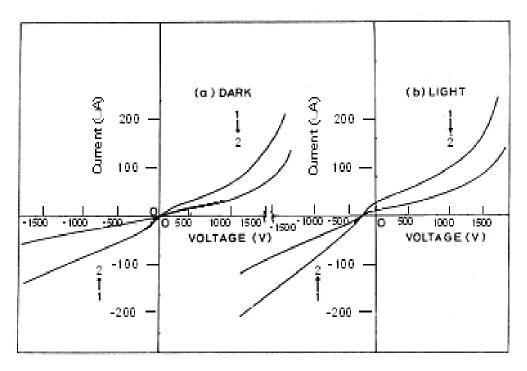


Fig.2. Linear plots of the current-voltage characteristics are shown for the cells in the dark (a) and with light (b) when the junctions are both forward and in reversed biased. The curves 1-2 represent Rose Bengal-TEA and Rose Bengal-Glucose respectively. The low resistivity of Rose Bengal-TEA compared to that of other dye-reductant films results in relatively higher currents at various voltages.

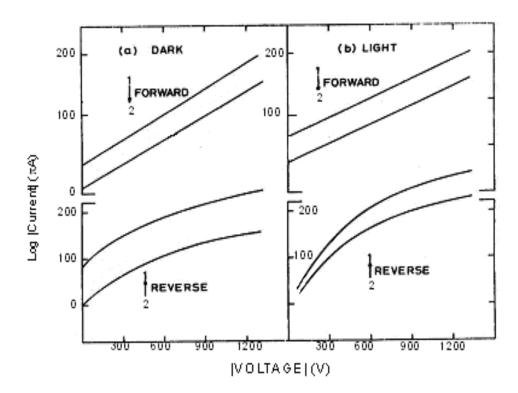


Fig.3. Semilog plots of current vs voltage for cells in the dark (a) and with light (b). The curves 1-2 represent Rose Bengal-TEA and Rose Bengal-Glucose respectively.