

Role of Ponceau-S-Mannitol System for Generation of Electrical Energy in Photogalvanic Cell

Mahesh Chandra^{1,*} and R. C. Meena²

1. Department of Chemistry, Deshbandhu College, New Delhi-110019

2. Department of Chemistry, Jai Narain Vyas University, Jodhpur (Rajasthan)-342 005

e-mail: dr.mahesh10@yahoo.co.in; rcmeena007@rediffmail.com

Abstract

*Ponceau-S is used as a photo sensitizer in photogalvanic cell for solar energy conversion and mannitol is used as an electron donor in this study. The effects of various parameters like pH, light intensity, diffusion length, reductant concentration and dye concentration on the electrical output of the cell is studied. The photopotential and photocurrent generated by this cell are 1080 mV and 240 μ A, respectively. The current voltage (*i*-*V*) characteristic of the cell is also studied and a mechanism for the generation of photocurrent is proposed.*

Keywords: Photogalvanic cell, Photopotential, Ponceau-S, Mannitol, Photocurrent

Introduction

The global warming and the rapid decrease in energy resources caused by the large-scale consumption of fossil fuels have become serious problem. Accordingly, renewable energy resources are attracting a great deal of attention and solar energy will be one of the most promising future energy resources. The photo effects in electrochemical systems were first reported by Becquerel^{1,2}. It has been reported that only negative photo potential should be obtained with carbonyl compounds³. Alonso et al. reported the use of electrodeposited CdSeO-5 TeO-5 electrode for solar energy conversion⁴. Jana and Bhowmik reported enhancement in the power output of a solar cell consisting of mixed dyes⁵. Hara et al. investigated design of new coumarin dyes having thiophene moieties for highly efficient organic dye-sensitized solar cells⁶. It has been reported the use of toluidine blue nitroloacetic acid (TB-NTA)⁷, in Azur A-Glucose⁸ and Fluoroscein-EDTA¹⁰ systems. Similarly it has been reported that the photo galvanic cells for classroom investigation and femto-second excited state dynamics of an iron (II) polypyridyl solar cell¹². Schwarzburg and Willig explored the origin of photovoltage and photocurrent in nanoporous, dye-sensitized, photo electrochemical solar cell¹³. The sensitization of nanoporous films on TiO₂ with santaline (red sandal wood pigment) and the construction of a dye-sensitized solid state photovoltaic cell was attempted by Tennakone and Kumara¹⁴. Yadav et al. reported use

*Corresponding author

of bismarck brown-ascorbic acid (BB-AA) system in photo galvanic cell for solar energy conversion¹⁵. Photo galvanic cells containing reductants and photo sensitizers were reported¹⁶⁻²⁰.

The research in the field of photogalvanic cells is still in its infancy with respect to its viability and practical applicability. Therefore, requires thorough exploration to increase the conversion efficiency and storage capacity by selecting a suitable redox couple and photo sensitizers. A detailed survey of literature revealed that no attention has been paid to the ponceau-S -mannitol system in a photo galvanic cell for solar energy conversion. In this context, the present work has been undertaken.

Experimental Methods

All the solutions were prepared in doubly-distilled water and stored in amber-colored containers to protect them from light. A mixture of the solution of the dye, mannitol, sodium hydroxide and water were filled into an H-shaped glass cell. A platinum electrode ($1 \times 1 \text{ cm}^2$) was placed in one compartment of the cell and a reference saturated calomel electrode (SCE) in the other compartment. The platinum electrode was exposed to a 200 W tungsten lamp while the SCE was kept in the dark. The temperature of the system was maintained at 303 K (± 0.1). A water filter was used to cut-off infrared radiations. A digital pH meter and a microammeter were used to measure the potential and current, respectively. The current-voltage characteristics were determined by applying extra load with the help of carbon pot ($\log 500 \text{ K}$) connected in the circuit. With this variable resistor (carbon pot) current-voltage curve was plotted.

Results and discussion

Effect of pH

The effect of pH on the electrical output of the cell is shown in Fig. 1. Photo potential and photocurrent are increased with increasing pH until at pH 13. Further increase in pH results in a decrease in the electrical output of the cell.

The dependence of photo potential and photocurrent on the concentration of the dye was studied and the results are shown in Fig. 2. On increasing the concentration of ponceau-S, both the photo potential and the photocurrent increase till a maximum is achieved at $4.8 \times 10^{-6} \text{ M}$, after which both characteristics are decreased. A small output is obtained at a low concentration of ponceau-S because a smaller number of dye molecules are available for excitation and consecutive donation of electrons to the platinum electrode. A large concentration of dye results in a decrease in photo potential because the intensity of light reaching the dye molecules (near the electrode) decreases due to the major portion of the light being absorbed by the dyes available in its path.

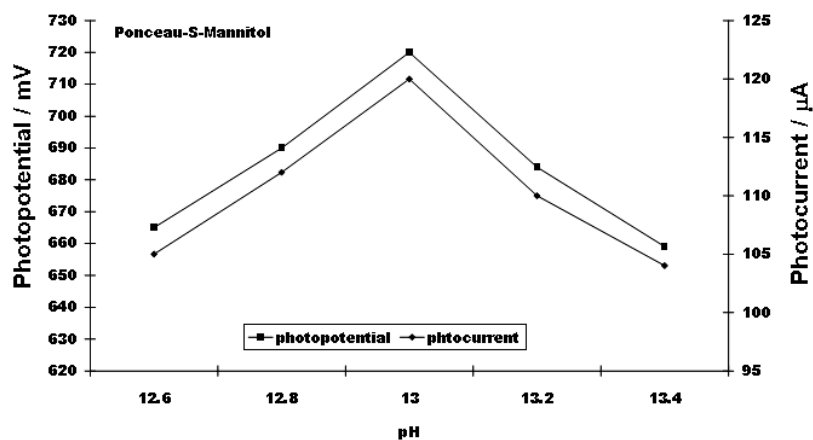


Figure 1: Variation of photopotential and photocurrent with pH Effect of ponceau-S concentration

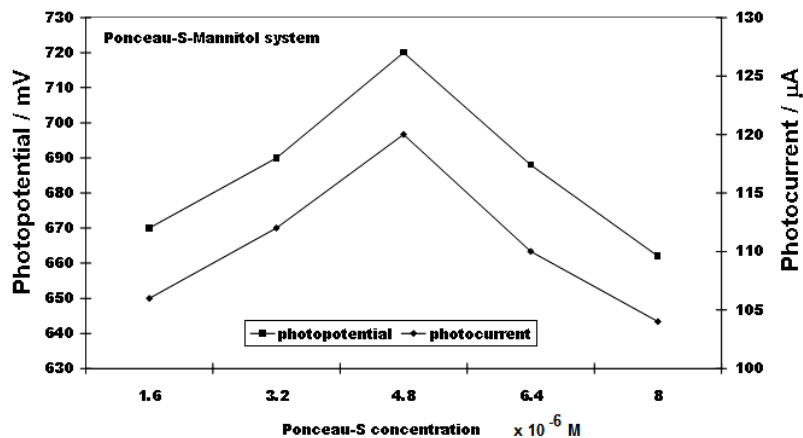


Figure 2: Variation of photopotential and photocurrent ponceau-S concentration effect of mannitol concentration

The dependence of photopotential and photocurrent on the concentration of the reductant (that is, mannitol) was studied and the results are shown in Fig. 3. Both the photopotential and the photocurrent achieve maximum values at the concentration of 2×10^{-3} M of mannitol. At low concentrations, the power output is small due to the fewer number of reductant molecules available for electron donation to the dye molecules, whereas a large concentration of reductant hinders the movement of dye molecules reaching the electrode in the desired time limit.

The variation of two electric parameters with light intensity is shown in Fig. 4. The photocurrent is linearly increased with increasing in the intensity of the light, whereas the photopotential is increased in a logarithmic manner. The number of photons per unit area (incident power) that strike the dye molecules around the platinum electrode increases with the increase in the light intensity. Hence, the photocurrent and the photopotential of the photo galvanic cell are favorably increased.

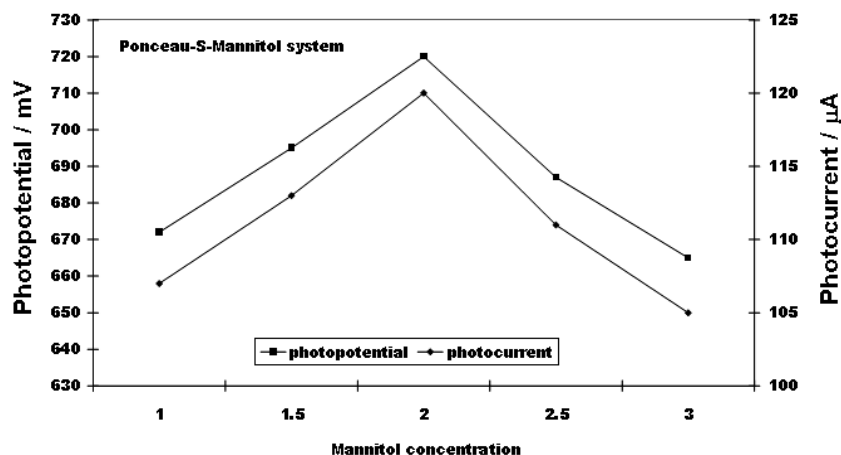


Figure 3: Variation of photopotential and photocurrent with mannitol concentration
Effect of light intensity

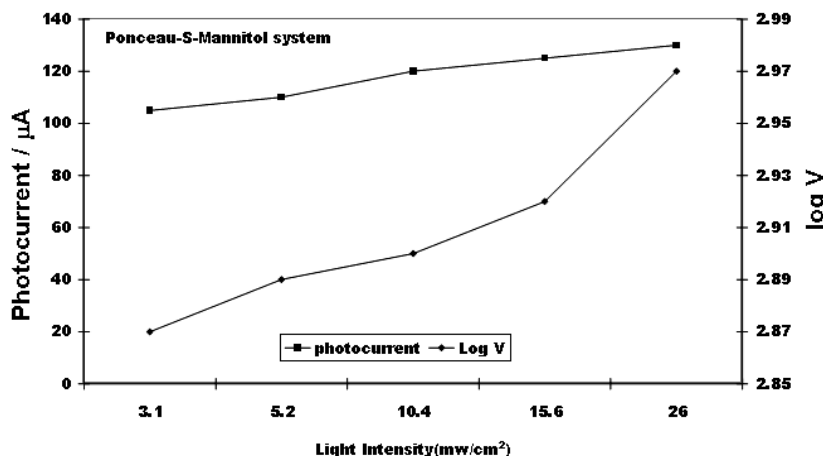


Figure 4: Variation of photopotential and photocurrent with light intensity
Effect of diffusion length

H-cells of different dimensions were used to study the effect of the variation of diffusion length on the current parameters of the cell (i_{max} , i_{eq} and initial rate of current generation). The results are shown in Fig. 5. There is a sharp increase in photocurrent (i_{max}) initially. This behavior indicates an initial rapid reaction, followed by a slow rate-determining step at a later stage.

Current voltage (i - V) characteristics, conversion efficiency and performance of the cell

The open-circuit voltage (V_{oc}) and short-circuit current (i_{sc}) of the photogalvanic cell were measured by means of a digital multi-meter (keeping the circuit open) and a micro-ammeter (keeping the circuit closed), respectively. The current and potential between two extreme values (V_{oc} and i_{sc}) were recorded with the assistance of a carbon pot (linear 470 K) that was connected in the circuit of the multi-meter and through which an external load was

applied. The i-V characteristic of the cell containing a ponceau-S-mannitol system is shown in Fig. 6. With the help of the i-V curve, the fill factor and conversion efficiency of the cell are found to be 0.45 and 0.4560 %, respectively, using the formula:

$$\text{Fill Factor} = \frac{V_{pp} \times i_{pp}}{V_{oc} \times i_{sc}}$$

$$\text{Conversion Efficiency} = \frac{V_{pp} \times i_{pp}}{10.4mWcm^{-2}} \times 100 \%$$

The potential and the current at the power point [A point in the i-V curve is called the power point (pp) and was determined where the product of photocurrent and photo potential is maximum] are represented by V_{pp} and i_{pp} , respectively.

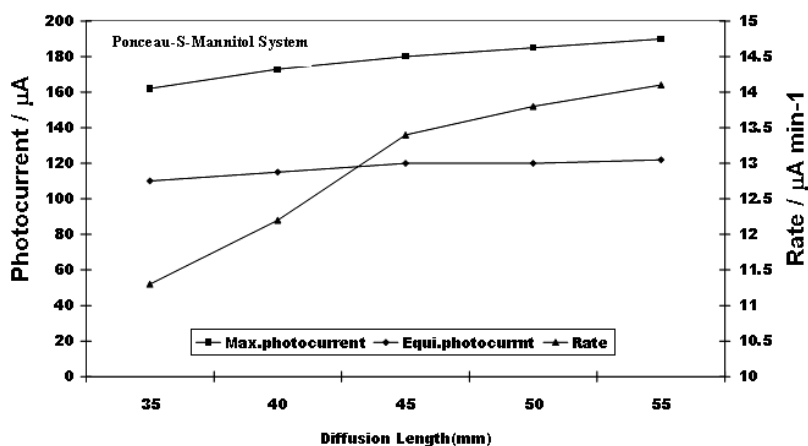


Figure 5 : Variation of current with diffusion length

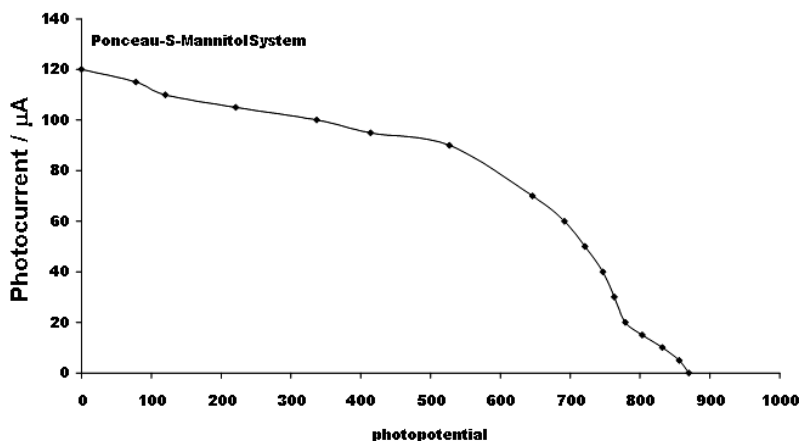


Figure 6: Current-potential (i-V curve) of the ponceau-S mannitol cell systems

The performance of the cell was studied by applying the external load that was necessary to have the current and the potential at the power point after removing the source

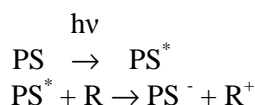
of light. The cell can be used in the dark at its power point for 36 minutes. Whereas photovoltaic cell cannot be used in the dark even for a second, a photogalvanic system has the advantage of being used in the dark but at lower conversion efficiency.

Mechanism

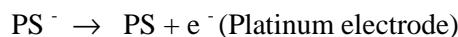
As no reaction is observed between the ponceau-S and mannitol in the dark, it may be concluded that the redox potential of mannitol is much higher than that of ponceau-S. A rapid fall in potential is observed when the platinum electrode is illuminated. The potential reaches a steady value after certain period of exposure. Although the direction of the change of potential is reversed on removing the source of light, the potential does not returns to its initial value. This means that the main reversible photochemical reaction is also accompanied by some side irreversible reactions. The electro active species in this photogalvanic system is thus different from that of the well-studied thionine–iron (II) system. In the present case, the leuco- or semi reduced dye is considered to be the electrode active species in the illuminated chamber and the dye itself in dark chamber. On the basis of the information gained above, the mechanism of photocurrent generation in the photogalvanic cell can be represented as follows:

Illuminated Chamber

Bulk solution

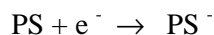


At electrode

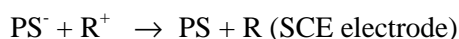


Dark chamber

At electrode



Bulk solution



where, R, R⁺, PS, PS⁻ are the reductant mannitol, its oxidized form, ponceau-S and its leuco or semileuco forms, respectively.

Conclusions

On the basis of above results, it is concluded that ponceau-S can be used successfully as a photosensitizer in a photogalvanic cell. The conversion efficiency of the cell is 0.4560% and the cell can be used in dark at its power point for 36 minutes. Photovoltaic cells have better conversion efficiency than photo galvanic cells. Photogalvanic cells have the

advantages of having in-built storage capacity. Thus, photogalvanic cells show good prospects of becoming commercially viable.

Acknowledgements

The Authors is grateful to Principal, Desh Bandhu College, Kalkaji, New Delhi for providing the necessary laboratory facilities to conduct this research works.

References

1. E. Becquerel, *C.R. Acad. Sci. Paris.*, 1839, **9**, 145
2. E. Becquerel, *C. R. Acad. Sci. Paris*, 1839, **9**, 561
3. J. J. Surash and D. M. Hercules, *J. Phys. Chem.*, 1962, **66**, 1602
4. V. N. Alanso, M. Belay, P. Chartier and V. Ern, *Rev. Phys. Appl.*, 1981, **16**, 5
5. A. K. Jana and B. B. Bhowmik, *J. Photochem. Photobiol.*, 1999, **122A**, 53
6. K. Hara, M. Kurashige, Y. Dan-oh, C. Kasada, A. Shinpo, S. Suga, K. Sayama and H. Arakawa, *New J. Chem.*, 2003, **27**, 783
7. S. C. Ameta, R. Ameta, S. Seth and T. D. Dubey, *Afinidad*, 1998, **XLV**, 264
8. S. C. Ameta , S. Khamesare , R. Ameta and M. Bala , *Int. J. Energy Res.*, 1999,**14**, 163
9. S. C. Ameta, P. B. Punjabi, J. Vardia, S.Madhwani and S.Chaudhary, *J. Power Sources*, 2006, **159**, 747
10. S. Madhwani, R. Ameta, J. Vardia, P. B. Punjabi and V. K. Sharma, *Energy Sources*, 2007, **29**, 721
11. C. Bohrmann-Linde and M. W. Tausch, *J. Chem. Educ.*, 2003, **80**, **1471**
12. J. E. Monat and J. K. McCusker, *J. Amer. Chem. Soc.*, 2000, **122**, 4092
13. K. Schwarzburg and F. Willig, *J. Phys. Chem.*, 1999, **103B**, 5743
14. K.Tennakone and R. A. Kumara GR, *J. Photochem. Photobiol.*, 1998, **117A**, 137
15. Sushil Yadav , R. D. Yadav and Gautam Singh , *Int. J. Chem. Sci.*, 2008, **6(4)**, 1960
16. R. C. Meena, Gautam Singh and K. M. Gangotri, *Afinidad*, 2003, **59 (501)**, 253
17. R. C. Meena and R.S. Sindal., *Int. J. Chem. Sci.*, 2004, **2(3)**, 321
18. Suresh C. Ameta, Sadhana Khamesra, Anil K. Chittoro and K. M. Gangotri., *Int. J. Energy Res.*,1989, **13**, 643
19. K. M. Gongotri, R. C. Meena and Rajni Meena., *J. Photochem. Photobiol. A: Chem.*, 1999, **123**, 93
20. R. C. Meena, *J. Indian Chem. Soc.*, 2008, **85**, 280