

# DESIGN AND BIOLOGICAL EVALUATION OF HYDROGEL BIOCOMPOSITES WITH ANTI-OBESITY ACTIVITY.

S.J. OWONUBI<sup>a</sup>, B. A. ADERIBIGBE<sup>b</sup> AND E. R. SADIKU<sup>a</sup>

<sup>a</sup> Department of Polymer Technology, Tshwane University of Technology, CSIR Campus, Building 14D, Private Bag X025, Lynwood Ridge 0040, Pretoria, South Africa. [oshesan@gmail.com](mailto:oshesan@gmail.com), [sadikur@tut.ac.za](mailto:sadikur@tut.ac.za)

<sup>b</sup> Department of Chemistry, University of Fort Hare, Alice Campus, Eastern Cape, South Africa. [blissingaderibigbe@gmail.com](mailto:blissingaderibigbe@gmail.com)

Type II diabetes is a chronic condition that is progressive with no effective cure till date. Glycaemic control is very critical so as to reduce long-term micro and macro-vascular complications resulting from its progressive nature. Hydrogel biocomposites prepared from a combination of thermally reduced graphene oxide, natural and synthetic polymers were encapsulated with anti-hyperglycemic drugs in selected ratios. The biocomposites were further characterized by Fourier Transform Infrared Spectroscopy (FTIR), Thermogravimetric Analysis (TGA), Scanning Electron Microscopy (SEM), Transmission Electron Microscopy (TEM) and RAMAN. Successful drug encapsulation was confirmed by Fourier transform Infrared and UV-vis spectroscopy. In vitro drug release studies were performed at selected physiological pH values. The hydrogels biocomposites exhibited controlled release profile at pH 7.4 and a sustained release profile at pH of 1.2 suggesting that the hydrogels are potential targeted drug delivery systems. Biological evaluation was performed against 3T3-L1 pre-adipocytes cell lines and successful expression of the gene regulators confirmed that their anti-obesity activities were not significantly different from positive control metformin.

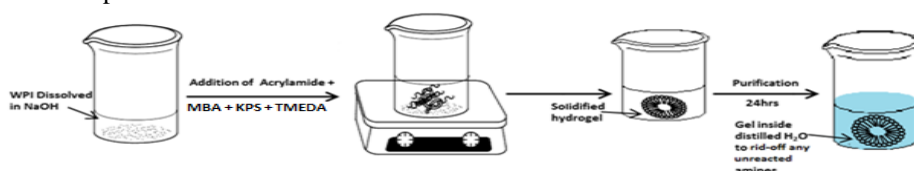


Fig. 1: Preparation of hydrogel by free radical polymerization

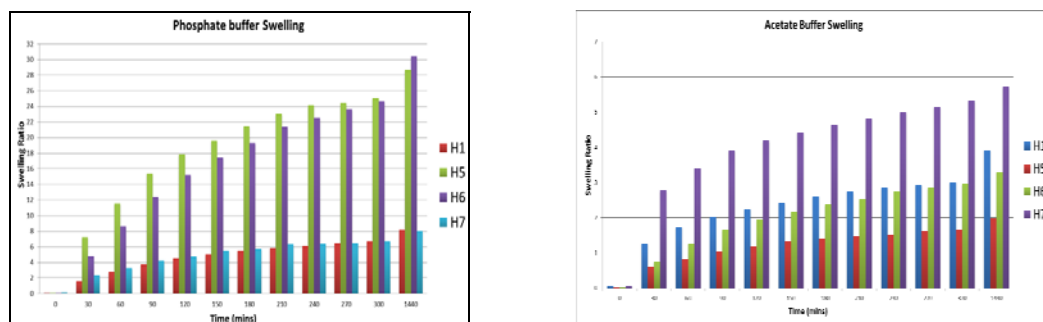


Fig. 2: Swelling rate of hydrogel biocomposite in buffer solutions.

- **Acknowledgement:** Prof E.R Sadiku research group, Prof. Emmanuel Mukwevho and the Diabetes research group, and DST/CSIR National Center for Nanostructured Materials, Pretoria, for their immense moral and technical support.

## References

- <sup>1</sup>Gudeman, L. F.; Peppas, N. A. *Journal of Applied Polymer Science*. **1995**, 55,919–928.
- <sup>2</sup>Gupta, P.; Vermani, K.; Garg, S. *Drug Discovery Today*. **2002**, 10, 569-579.
- <sup>3</sup>Peppas, N.A.; Buresa, P.; Leobandunga, W.; Ichikawab, H. *European Journal of Pharmaceutics and Biopharmaceutics*. **2000**, 50, 27-46.