



627/E2/Poster

A model system to investigate drug loading and releasing using bacterial nanocellulose and bovine serum albumin

W. R. A. P. J. Ratnayake,^{1,5} J.W. Damunupola,² A. L. C. J. Liyanage,³ R. G. S. C. Rajapaksha,⁴
and A. C. A. Jayasundera^{5*}

¹Postgraduate Institute of Science, University of Peradeniya, Peradeniya

²Department of Botany, Faculty of Science, University of Peradeniya, Peradeniya

³Department of Food Science & Technology, Faculty of Applied Sciences, Sabaragamuwa University of Sri Lanka, Belihuloya

⁴Department of Molecular Biology & Biotechnology, Faculty of Science, University of Peradeniya, Peradeniya

⁵Department of Chemistry, Faculty of Science, University of Peradeniya, Peradeniya

In the present study, the applicability of freeze dried bacterial nanocellulose (fd-BNC) synthesized by *Gluconacetobacter* spp. as a drug delivery matrix for proteins using Bovine Serum Albumin (BSA) as a model drug was systematically investigated. BSA was chosen because of its high-water solubility, abundance, good stability, and wide acceptance. BNC possesses superior mechanical (high tensile strength), chemical (hydrophilicity, high degree of polymerization, crystallization), and biological (non-cytotoxic, non-biodegradable, high purity) properties. The study focused on investigating different drug concentrations of BSA—2, 5, 10, 20 (mg/ml)—dissolved in phosphate buffered saline (PBS) adsorbed and released from pre-swelled BNC hydrogel at predetermined time points under shaking conditions, at room temperature. The loading profile, determined by UV absorbance spectrometry at 278 nm, demonstrated mean loaded BSA amounts of 1.65 (\pm 0.11), 3.02 (\pm 0.41), 7.04 (\pm 0.66), and 13.52 (\pm 0.53) (mg) after 48 hours for 2, 5, 10, and 20 mg/ml, respectively. The regression coefficient (r^2) of the equilibrium loading isotherm was 0.9965. Higher BSA adsorption may be due to the fine network of nanofiber structure in BNC. Drug loading stabilized after the initial 8 hours, and was controlled mainly by diffusion, whereas no saturation effects were observed. The release profile indicated a 'burst release' during the initial 8 hours, followed by a controlled release. The mean released amount of 2, 5, 10, and 20 (mg/ml) BSA concentrations were 1.12 (\pm 0.19), 2.89 (\pm 0.27), 5.59 (\pm 0.44) and 10.37 (\pm 0.53) (mg) respectively. The release stabilized after 8 hours, with a constant amount of the drug being released. Diffusion exponent (n) values were 0.65, 0.67, 0.79, and 0.69 for 2, 5, 10, and 20 (mg/ml) BSA. The Ritger–Peppas Power law model suggested that the studied BSA-fd-BNC system's drug transport mechanism is Non-Fickian (anomalous), an overlay of diffusion and swelling controlled release being observed. The present study suggests BNC as an innovative and attractive biopolymer for controlled drug delivery.

Keywords: Bacterial nanocellulose, controlled release, bovine serum albumin, anomalous diffusion

E-mail: acaj@pdn.ac.lk