

INTERACTION OF CARBON NANOTUBE AND CELLULOSE NANOFIBER WITH ALGAL CELLS *KLEBSORMIDIUM FLACCIDUM*

Michele M Pereira¹, Ludovic Mouton², Claude Yéprémian³, Alain Couté³, Joanne Lo⁴, José M Marconcini⁵, Luiz O Ladeira⁶, Nádia RB Raposo¹, Humberto M Brandão⁷ and Roberta Brayner², (1) UFJF – s/n, rue José Lourenço Kelmer 36036-900 Juiz de Fora / Brazil (2) ITODYS – 15, rue Jean-Antoine de Baïf 7086 Paris / France (3) Muséum National d'Histoire Naturelle – 57, rue Curvier 75005 Paris / France (4) Institut Jacques Monod – 15, rue Hélène Brion 75013 Paris / France (5) Embrapa Instrumentation – 1452, rue XV de Novembro 13560-970 São Carlos / Brazil (6) UFMG – 6627, rue Presidente Antonio Carlos 31270-901 Belo Horizonte / Brazil (7) Embrapa Dairy Cattle – 610, rue Eugenio do Nascimento 36038-330 Juiz de Fora / Brazil.

Multi-walled carbon nanotubes (MWCNTs) and cellulose nanofibers (CNF) are noteworthy nanoparticles (NPs), which encompass a number of potential applications, being used in water treatment, cosmetics, as well as reinforcement materials, biosensors and medical equipment. However, with the rise of nanotechnologies, the risk of contamination of aquatic ecosystems with NPs is increasing. Thus, the aim of this study was to evaluate the MWCNT and cotton CNF toxicological effects on freshwater green microalgae *Klebsormidium flaccidum*. *K. flaccidum* was grown in sterile *Bold's Basal* (BB) culture medium at pH 7.4 at a controlled temperature of $20.0 \pm 0.5^\circ\text{C}$ and luminosity of $50\text{--}80 \mu\text{mol m}^{-2} \text{s}^{-1}$ photosynthetic photon flux. Appropriate concentrations of each nanomaterial stock solution ($1, 50$ and $100 \mu\text{g ml}^{-1}$) were added to a microalgal culture in the exponential growth phase and incubated for 24, 48, 72 and 96 hrs. Cell viability was measured by a trypan blue dye exclusion test. In order to evaluate morphological, cellular ultrastructure changes and interaction between NPs and *K. flaccidum*, we analyzed microalgae cells by Scanning electron microscopy (SEM) after 48 h of contact with MWCNT and cotton CNF ($100 \mu\text{g ml}^{-1}$). Data were analyzed by ANOVA and differences among means were compared by the Student–Newman–Keuls' test using the general linear model by SAS version 9.1. Differences between different groups were considered statistically significant at $P < 0.05$. NPs significantly decreased cell viability ($P < 0.05$), depending on concentration and time (Fig. 1). The cell shrinkage was noted on the cells treated with both MWCNTs and cotton CNFs (Fig. 2B and 2C). In conclusion, we have demonstrated that exposure to MWCNTs and to cotton CNFs affects cell viability and algal cell morphology.

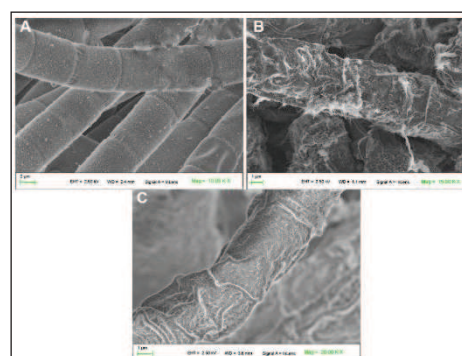
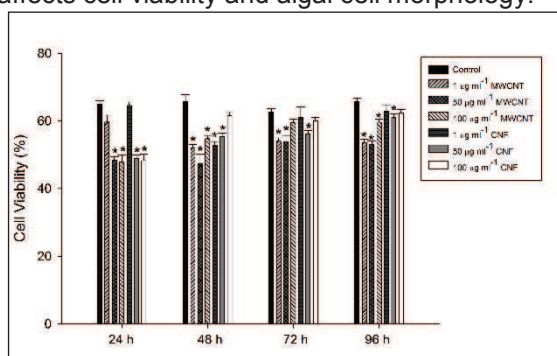


Fig. 1 – Effect of MWCNTs or CNF on cytotoxicity of *K. flaccidum* in vitro cultured in BB medium. *Asterisks denote a significant difference from the control group. Calculated probability (* $P < 0.05$).

Fig. 2 – SEM images of *K. flaccidum* exposed to $100 \mu\text{g mL}^{-1}$ MWCNTs or cotton CNF for 48h in BB medium. A: Control; B: MWCNTs; C: CNF.

Acknowledgments: This work was supported by CNPq, CAPES and Rede AgroNano.