SYNTHESIS OF DENDRIMER-CARBON NANOTUBE CONJUGATES

<u>García A.</u>^{##}, <u>Herrero M.A</u>.[†], Frein S.[§], Deschenaux R.[§], Muñoz R.[#], Bustero I.[#], Prato M.[†] [†]Dipartamento de Scienze Farmaceutiche, Università di Trieste, Piazzale Europa 1, 34127 Trieste (Italy) [#]Department of Biomaterials and Nanotechnologies, Inasmet Foundation, Mikeletegi Pasealekua 1, Teknologi Parkea, San Sebastián (Spain) [§]Institut de Chimie, Université de Neuchâtel, Av. de Bellevaux 51, Case postale 158, 2009 Neuchâtel, Switzerland

ainara.garcia@inasmet.es

Carbon Nanotubes (CNTs) have interesting physical and chemical properties that open attractive possibilities in many research areas. The main problem for further applications is the chemical inertness of the CNTs that makes difficult the attachment of molecules at the surface. The objective of our work is a high degree of functionalization on CNTs with a covalent attachment of the interesting molecules without provoking damage to the conjugated π -system. One approach is the attachment of dendrimers at the SWNTs with high density of functional groups that serve as anchor point for further reactions. For that, we have carried out a primary modification with 1,3 dipolar cycloadition reaction [1] with different kinds of commercial CNTs. Once the terminal Boc groups were deprotected and the amount of functionalities was quantified in each sample, the most reactive CNTs were chosen. Finally, dendrimers attachments were performed by carbodiimide chemistry and in all cases the final products were fully characterized.



Figure 1. AFM image of SWNT modified with the cycloadition reaction and diameter distribution of the sample (top) and that of the SWNT- dendrimer conjugated (down).

References:

[1] (a) Georgakilas, V., Kordatos, K., Prato, M., Guldi, D.M., Holzinger, M., Hirsch, A., J. *Am. Chem. Soc.*, **124** (2002) 760. (b) Georgakilas, V., Tagmatarchis, N., Pantarotto, D., Bianco, A., Briand, J.–P., Prato, M., *Chem. Commun*, **24** (2002), 3050