

**SiO<sub>2</sub> nanoparticle-neuron interaction: activation of ionic channels and calcium influx**

Carla Distasi  
University of Piemonte Orientale  
Department of Pharmaceutical Sciences  
Novara Italy

SiO<sub>2</sub> nanoparticles (SiO<sub>2</sub> NPs) are promising tools in the field of nanomedicine. In this perspective, the knowledge of the mechanisms of interaction between the SiO<sub>2</sub> NPs and their targets is a prerequisite to the rational design of safe and efficient nanotools for laboratory and clinical applications. In previous works, we showed that non-toxic doses of 50 nm SiO<sub>2</sub> NPs induce strong and long lasting calcium influx together with membrane potential depolarization and modulation of the electrical activity in GT1-7 neuroendocrine cells. To clarify the mechanisms of interaction between SiO<sub>2</sub> nanoparticles and the neuronal plasmamembrane and to obtain a detailed biophysical characterization of the multiple pathways activated, we have combined calcium imaging and electrophysiological patch clamp techniques with a pharmacological approach. TRPV4, Connexins and Pannexin-like channels are the major components of inward currents elicited by the NPs. Furthermore, pre-incubation with the antioxidant N-acetyl-L-cysteine (NAC) strongly reduces the [Ca<sup>2+</sup>]<sub>i</sub> increase. These findings suggest that SiO<sub>2</sub> NPs directly activate a complex set of calcium-permeable channels, possibly by catalyzing free radicals production.