



Online LTP Seminar

Lecture 3

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Title: How can we exploit biomaterials for the benefit of plasma medicine?

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Abstract: Biomaterials are employed for tissue and organs regeneration or functional repair, including delivery of therapeutics. In bone regeneration and repair, incorporation of drugs to biomaterials has been investigated as a means of providing additional functionalities to the material, and plasma processes contributed to bone biomaterials ie. through polymerisation processes to modulate the drug release¹.

With the evolution of plasma devices, great advances have been made in therapies based in cold atmospheric plasmas (CAP)^{2,3}. CAP generates reactive oxygen and nitrogen species (RONS) which can be transferred to liquids and have shown to have selective anticancer effects towards osteosarcoma. Osteosarcoma (OS) is the most common primary bone cancer, presenting poor prognosis and difficult treatment. We have recently shown that CAP-treated Ringer's saline produced cytotoxic effects in human OS cell lines, while sustained viability was observed in healthy cells. Higher levels of DNA damage were found in OS cells leading to apoptotic cell death. This was confirmed in mouse OS tumor sections in organotypic culture⁴.

However, injection of a liquid in the body results in fast diffusion due to extracellular fluids and blood flow. Therefore, the development of efficient vehicles which allow local confinement and delivery of RONS to the diseased site is a fundamental requirement. In this sense, biocompatible polymers with ability to form 3D networks can be an alternative to deliver the plasma-generated RONS locally. We will discuss the generation of RONS (H_2O_2 , NO_2^- , short-lived RONS) in different hydrogels⁵, whether these modify the physic-chemical properties of the material⁶, and the biological effects associated to them.

1. K. Khurana et al. *Eur. Polymer Journal* 107, 25 (2018); 2. D. Graves, *Plasma Process*, 11, 1120 (2014); 3. A. Dubuc et al. *Ther Adv Med Oncol* 10, 1 (2018); 4. M. Mateu-Sanz et al. *Cancers*, 12(1), 227 (2020); 5. C. Labay et al., *Scientific Reports* 9, 16160 (2019); 6. I. Hamouda et al. *Polymer*, 192, 122308 (2020).

