CeO₂ nanoparticles as safe CT X-ray contrast imaging agents

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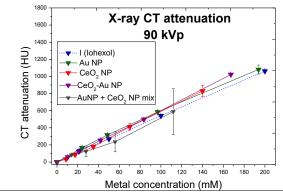
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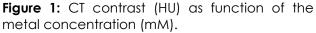
Use of X-ray computed tomography (CT) has increased dramatically over the last years in many countries owing to its availability, low cost and high resolution. An estimated 82 million scans were performed in the U.S. in 2016. However, a main concern about CT scans is the exposure to ionizing radiation. One study estimated that as many as 0.4% of current cancers in the U.S. are due to CTs performed in the past. Thus, an important issue is how to reduce the radiation dose during CT examinations without compromising the image quality. Contrast agents are regularly administered in order to improve image contrast. High Zbased nanoparticles have gained attention because they show notable lately advantages over commercial iodine-based compounds. For instance, they allow longer circulation times and deliver a large amount of heavy atoms[1]. In this work we present CeO₂ NPs as potential CT contrast agents. Cerium possesses high atomic number and electron density, it is biocompatible and CeO₂ NPs show antioxidant and antiinflammatory properties, providing radioprotective capabilities [2]. In addition, given that the contrast enhancement depends both on the tube potential (kVp) and energy spectrum, Au-CeO₂ hybrides have also been prepared, increasing thus Xray energy absorption range.

Firstly, it has been assessed the stability of the colloidal samples at the high

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concentrations required by the CT instruments (>100 mM), which are prone to aggregation and precipitation. In vitro CT results (Figure 1) indicate that, at equal concentrations, Ce-based molar NPs produce higher X-ray contrast than iodine. Additionally, we have observed that CeO₂ and Au-CeO2 NPs act as a free-radical scavenger (Figure 2), reducing thus the radiation adverse effects of using metals in CT imaging [3].





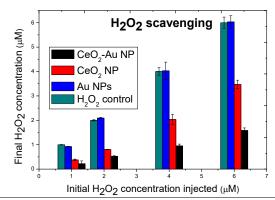


Figure 2: H₂O₂ scavenging tested with Amplex Red Assay

References

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- [2] Cafun JD et al. ACS Nano, 7 (2013)10726-32.
- [3] McMahon SJ et al. Scientific Reports 1:18 (2011) 1-8.