Dehydropeptide-based self-assembled hydrogels with incorporated Gd³⁺ chelates: potential Contrast Agents for MRI?

Teresa Pereira^a, Juan Gallo^b, Manuel Bañobre-López^b, Loic Hilliou^c, Paula M.T. Ferreira^a, and José A. Martins^a

^aCentre of Chemistry, University of Minho, Campus de Gualtar, 4710-057 Braga, Portugal; jmartins@química.uminho.pt

^b International Iberian Nanotechnology Laboratory (INL), Av. Mestre José Veiga s/n, 4715-330 Braga, Portugal

^cInstitute for Polymers and Composites, Department of Polymer Engineering, University of Minho, Campus de Azurém, 4800-058 Guimarães, Portugal

Magnetic Resonance Imaging (MRI) is at the forefront of clinical imaging. Paramagnetic relaxers (Gd³⁺ and Mn²⁺ chelates, iron oxide nanoparticles) shorten the relaxation times $(T_{1,2})$ *i.e.* enhance the relaxation rates $(R_{1,2}=1/T_{1,2})$ of the water protons in their vicinity yielding significant contrast enhancements-contrast agents. Supramolecular (selfassembled) hydrogels based on low molecular weight peptides are the new paradigm biomaterials: porous soft biocompatible materials made of highly hydrated fibrous 3D nanostructures, reminiscent of the extracellular matrix. Our research group developed selfassembled hydrogels based on dehydrodipeptides N-capped with naproxen (Npx, a NSAID drug). Dehydropeptide-based hydrogels exhibit resistance to proteolysis, are biocompatible and suitable nanocarriers for delivery of incorporated drugs. Recently, we demonstrated that incorporation of SPION endows dehydropeptide-based hydrogels with magnetic properties (magnetogels): hyperthermia and MRI reporting properties. In this communication we report novel supramolecular hydrogels prepared by co-assembly of dehydropeptides and Gd³⁺ chelates. The hydrogels are characterised regarding coassembly (fluorescence, CD spectroscopy) and micro- nano-structure (STEM) and rehologial properties. The co-assembled hydrogels are characterized also as Contrast Agents for MRI by ¹H relaxometry (60 MHz, 37°C) and MRI (120 MHz, 37 °C).

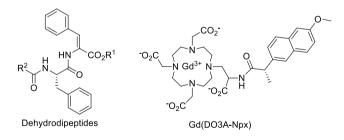


Figure 1

The authors acknowledge the FCT under the Project PTDC/QUI-QOR/29015/2017 and CQ/UM UID/QUI/00686/2013 and UID/QUI/0686/2016 projects.

M. F. Ferreira *et al.*, *Dalton Trans.*, **2014**, 43, 3162.
H. Vilaça *et al.*, *Biomacromolecules*, **2015**, 16, 3562.
Carvalho *et al.*, *Nanomaterials*, **2019**, 9, 541.