

Název: Postupy a přístupy pro syntézu
magnetizovatelných částic

Školitel: Doc. RNDr. Pavel Kopel, Ph.D.

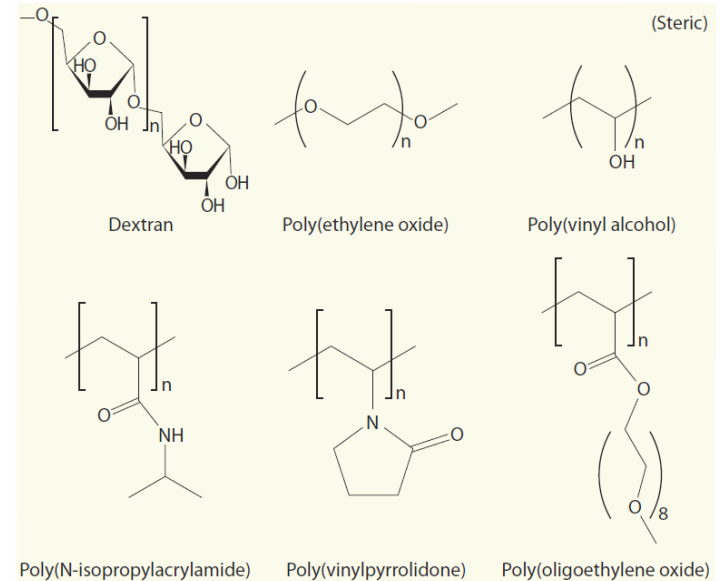
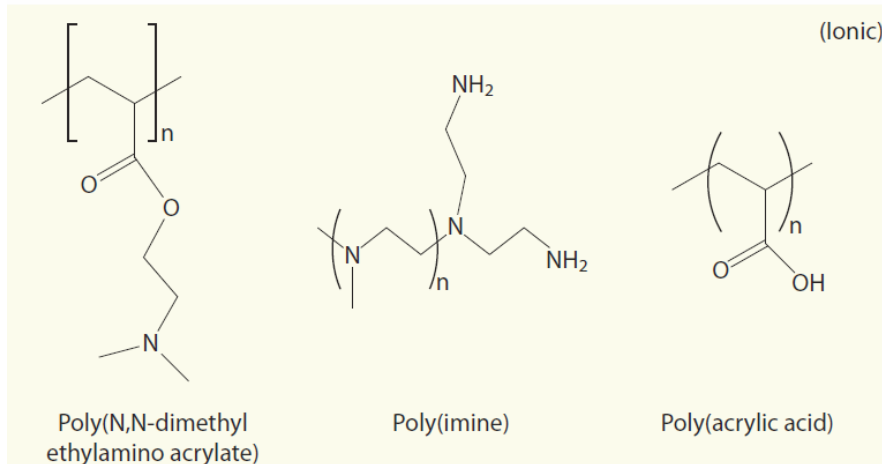
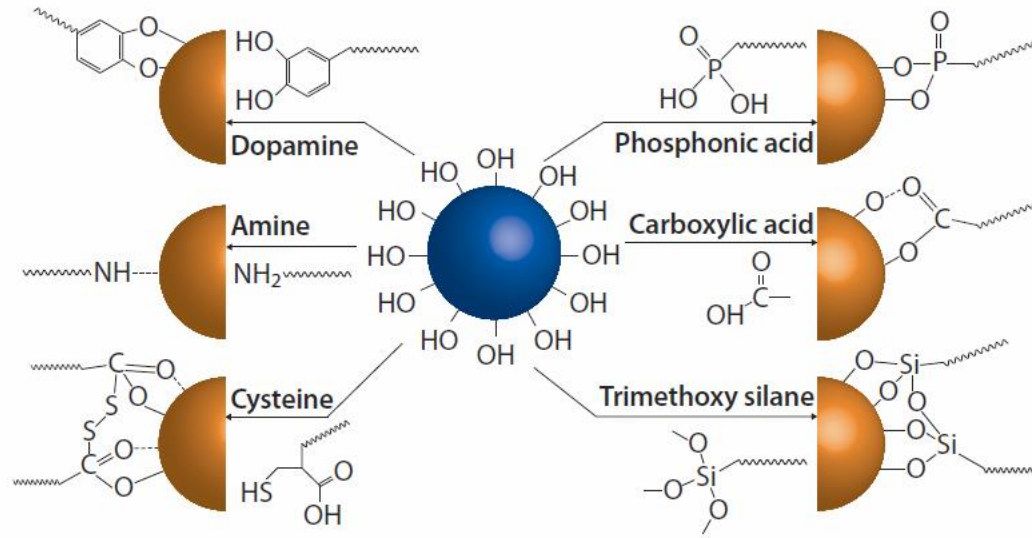
Datum: 6.11.2013

Synthetic methods for iron oxides preparation

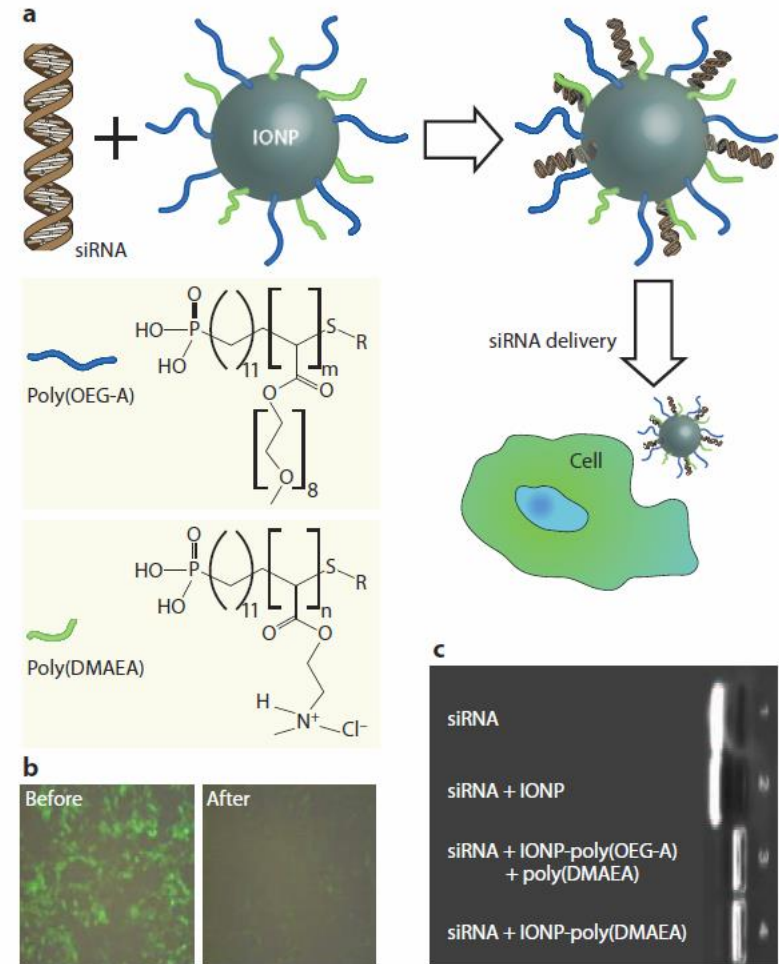
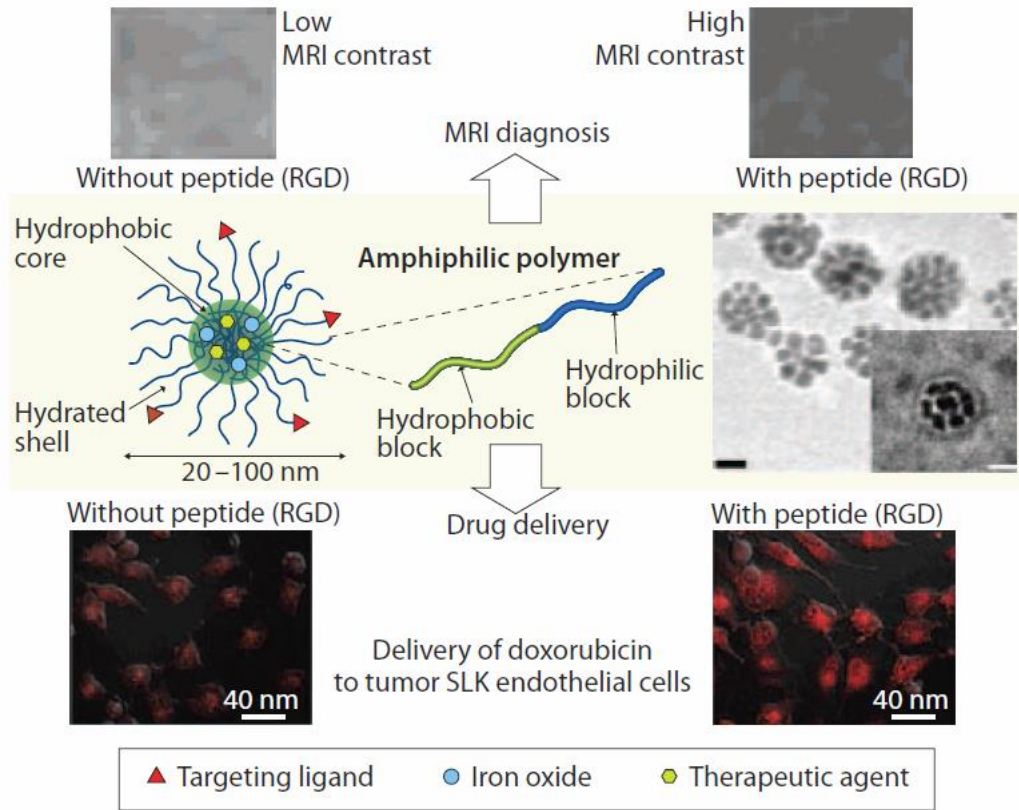
Synthetic method	Synthesis	Reaction temp. [°C]	Reaction period	Solvent	Surface-capping agents	Size distribution	Shape control	Yield
co-precipitation	very simple, ambient conditions	20–90	minutes	water	needed, added during or after reaction	relatively narrow	not good	high/ scalable
thermal decomposition	complicated, inert atmosphere	100–320	hours–days	organic compound	needed, added during reaction	very narrow	very good	high/ scalable
microemulsion	complicated, ambient conditions	20–50	hours	organic compound	needed, added during reaction	relatively narrow	good	low
hydrothermal synthesis	simple, high pressure	220	hours ca. days	water-ethanol	needed, added during reaction	very narrow	very good	medium

- Co-precipitation is a facile and convenient way to synthesize iron oxides (either Fe₃O₄ or γ-Fe₂O₃) from aqueous Fe²⁺/Fe³⁺ salt solutions by the addition of a base under inert atmosphere at room temperature or at elevated temperature. The size, shape, and composition of the magnetic nanoparticles depends on the type of salts used (e.g. chlorides, sulfates, nitrates), the Fe²⁺/Fe³⁺ ratio, the reaction temperature, the pH value and ionic strength of the media.
- Preparation of paramagnetic microparticles
- Maghemite nanoparticles were prepared by sodium borohydride (NaBH₄) reduction of iron chloride (FeCl₃·6H₂O) according to Magro et al. and Pucek et al [11, 14]. 10 g of FeCl₃ • 6H₂O was dissolved in 800 mL of water. Briefly, 2 g of FeCl₃•6H₂O was dissolved in 80 mL of MiliQ water and a solution of 0.2 g of NaBH₄ in ammonia (10 mL, 3.5 % m/v) was poured into the first solution with vigorous stirring. The obtained solution was heated at boiling temperature for 2 h. After cooling and standing for 2 h at room temperature, the obtained magnetic nanoparticles were separated by external magnetic field and washed several times with water. The nanoparticles, prepared as described above were suspended in 80 mL of water and Ti(isopropox)₄ was added. Resulting product was stirred on Biosan OS-10 through whole night. Then microparticles were separated by external magnetic field, washed with MiliQ water and finally dried at 40 °C.

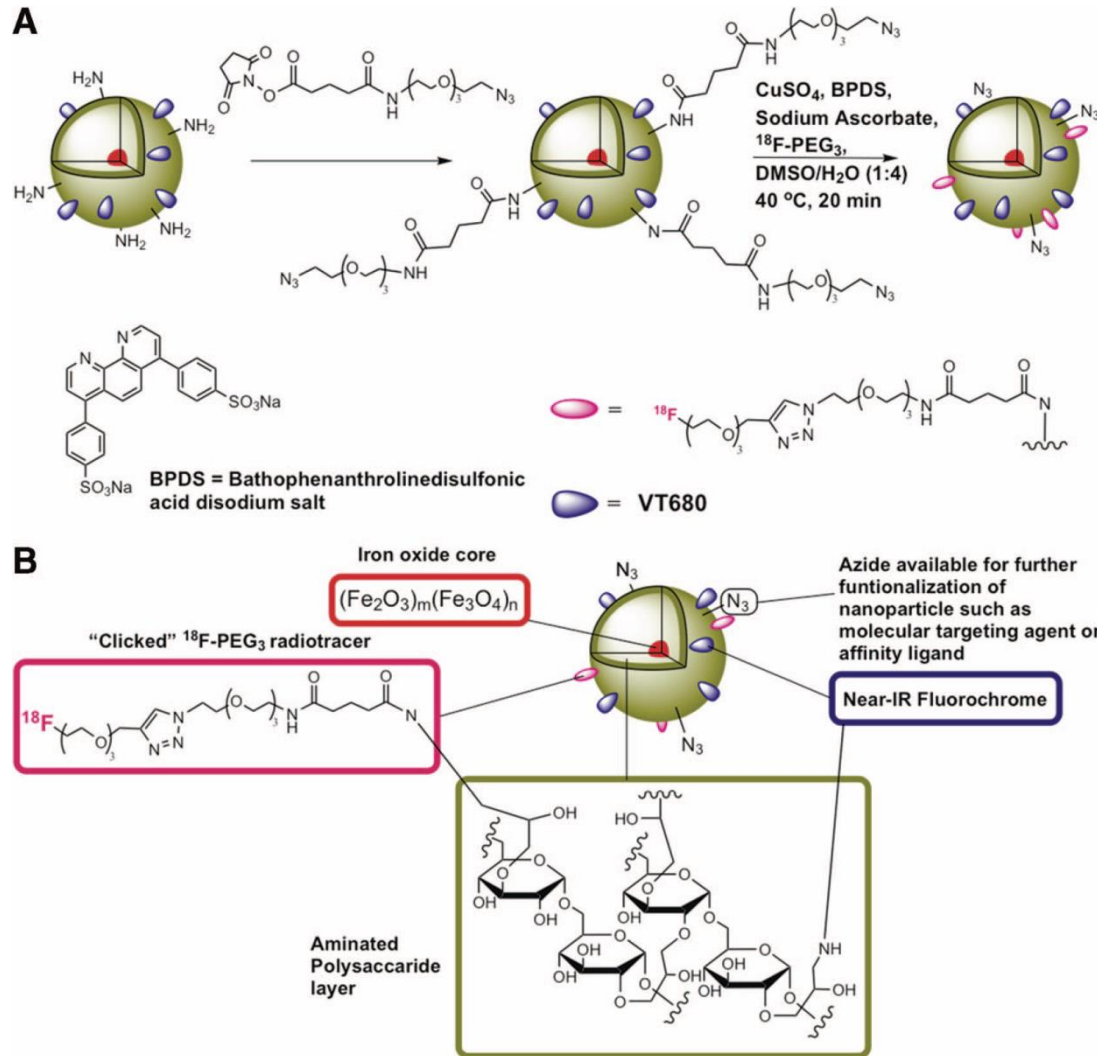
Coating of SPION nanoparticles



Delivery of drugs and siRNA

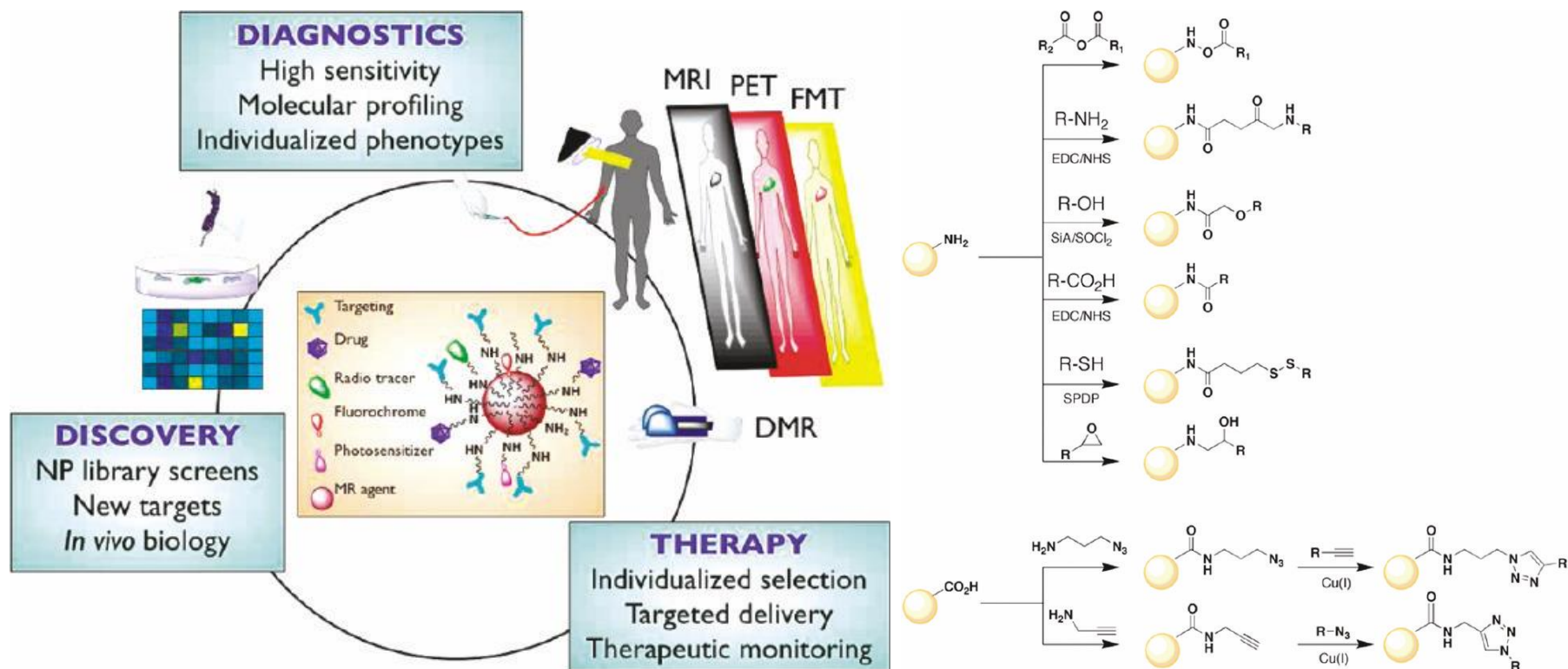


18F Labeled Nanoparticles for in Vivo PET-CT Imaging



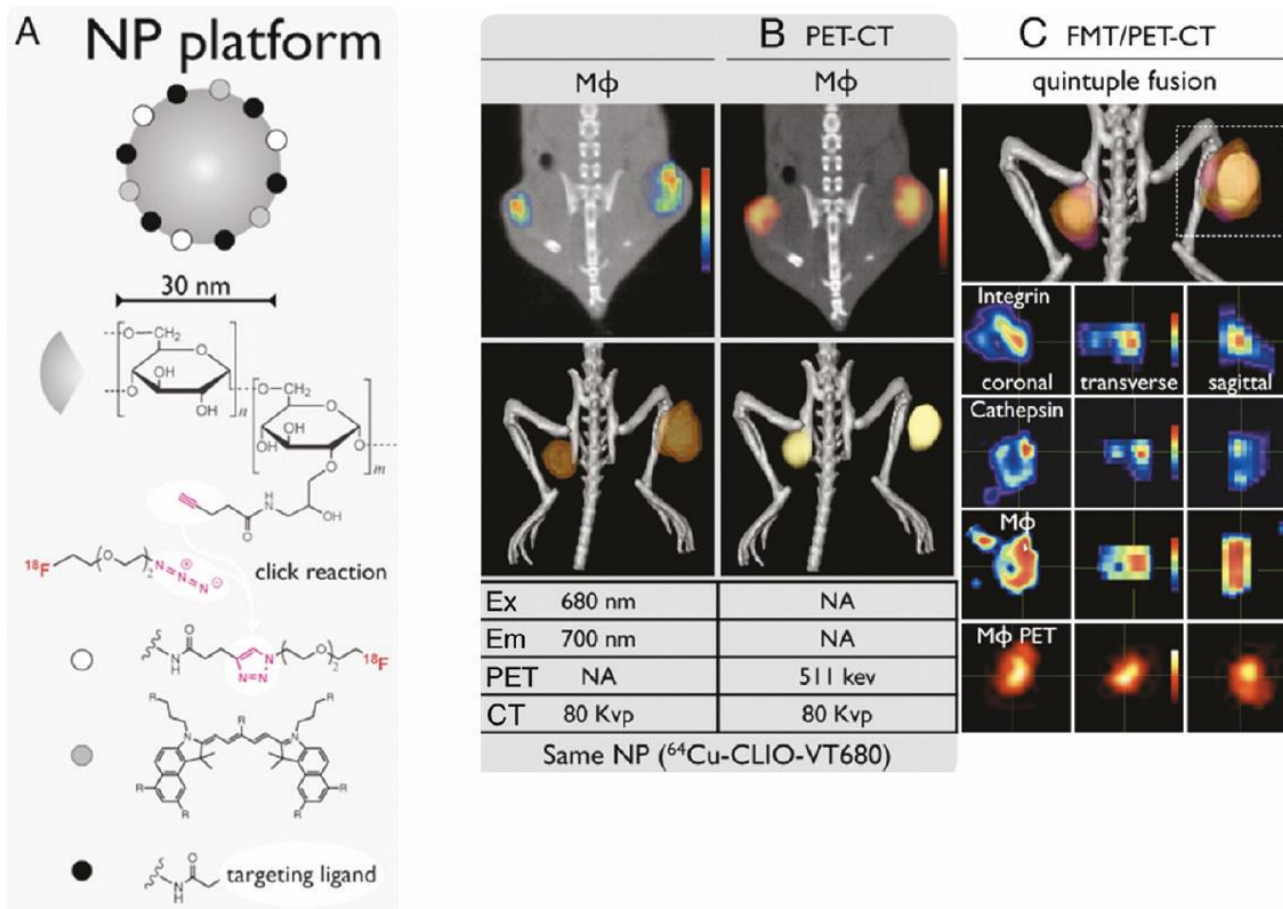
- Preparation of ^{18}F -CLIO. (A) Derivatization of primary amines on CLIO-VT680 (near-infrared fluorochrome Vivotag-680 (VT680)) with the NHS ester (N-Hydroxysuccinimide) of 1-azido-13-oxo-3,6,9-trioxa-12-azaheptadecan-17-oic acid followed by chemoselective "click" of ^{18}F -PEG₃ radiotracer. (B) Schematic of ^{18}F -CLIO.
- Bioconjugate Chem. 2009, 20, 397–401

Dextran-Coated Iron Oxide Nanoparticles: A Versatile Platform for Targeted Molecular Imaging, Molecular Diagnostics, and Therapy



- Conjugation chemistries to attach small molecules to CLIO
- ACCOUNTS OF CHEMICAL RESEARCH ' 842–852 ' 2011 ' Vol. 44, No. 10

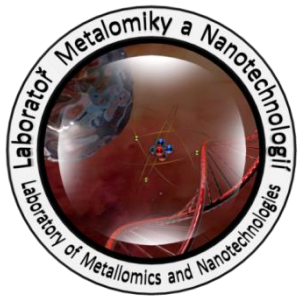
Dextran-Coated Iron Oxide Nanoparticles: A Versatile Platform for Targeted Molecular Imaging, Molecular Diagnostics, and Therapy



- Multimodal PET imaging using nanoparticles. (A) Versatile conjugation capabilities of CLIO, e.g., to ¹⁸F using click chemistry, but also to peptides or other targeting ligands. (B, C) In vivo multichannel PET-CT (B) and FMT/PET-CT (C) of tumor-bearing mice, coinjected with fluorescent peptide against integrins, a fluorescent cathepsin sensor, and ⁶⁴Cu-CLIO-VT680 (labeling macrophages).

Acknowledgements

All the members of Laboratory of Metalomics and Nanotechnology



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Thank you for Your attention

Reg.č.projektu: CZ.1.07/2.4.00/31.0023

Název projektu: Partnerská síť centra excelentního bionanotechnologického výzkumu