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Radio-labeled nano-platforms for imaging, image-guided drug delivery and theranostics

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The molecular imaging and nanotechnology laboratory at the University of Wisconsin - Madison is mainly focused on three areas: 1) development of multimodality molecular imaging agents; 2) nanotechnology and its biomedical applications; and 3) molecular therapy of cancer. In this talk, I will present our recent work on molecular imaging and image-guided drug delivery in cancer with a variety of nanomaterials. The primary imaging techniques used in this study are positron emission tomography (PET), photo-acoustic tomography (PAT), optical imaging and magnetic resonance imaging (MRI). Three of the major molecular targets that we are investigating are CD105 (i.e. endoglin), VEGFR and integrin $\alpha v \beta 3$. The nanomaterials that will be discussed in this presentation include nano-graphene oxide, zinc oxide nanomaterials, micelles, silica-based nanoparticles, magnetic nanoparticles, hybrid nanomaterials, among many others. A few representative side projects will also be presented, such as the facile synthesis of PET/MRI and PET/PAT dual-modality imaging agents.

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OncoCiDia: A Novel Depersonalized Combination of Targeted Chemo- and Radio-Theragnostics for the Management of Solid Malignancies

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Imaging and contrast media research has enabled discovery of small molecular necrosis-avid compounds (NACs) for the diagnosis of myocardial infarction and therapeutic assessment of tumor ablation with MRI, nuclear scintigraphy and optical imaging^{1,2}. The *in vivo* affinity of NACs to necrosis appears orders of magnitude higher than antigen-antibody, ligand-receptor and biotin-avidin interactions *in vivo*. Based on a soil-to-seeds hypothesis³, this stroma targetability is extended from diagnostic to theragnostic utilities by combined use of vascular disrupting agents (VDAs) such as CA4P to formulate a novel pan-anticancer approach, namely a small-molecular sequential dual targeting theragnostic strategy³. The dual targeting properties and conjugated iodine-131 that emits both beta and gamma radiations provide solid cancers (Onco) with both tumoricidal (Ci) and imaging diagnostic (Dia) effects, hence an acronym OncoCiDia³. Instead of directly attacking multmutant and refractory cancer cells (seeds) as in other cancer therapies, OncoCiDia selectively destroys and radioactively sterilizes the tumor microenvironment (soil)⁴. Multicenter preclinical investigations on the efficacy, safety, formulations and dosimetry suggest that this novel and unconventional anticancer strategy may present a relatively simple, workable, affordable and generic solution for diverse cancer problems, and deserve further exploitation.

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