## Time-resolved imaging and sensing in the near-infrared: Towards quantitative in vivo imaging

Yuyang Gu\*

University of Technology Sydney, Ultimo NSW 2007, Australia Institute for Biomedical Materials and Devices (IBMD) Ultimo NSW 2007, Australia \*Yuyang.Gu@uts.edu.au

For years, luminescence lifetime imaging has served as an indispensable tool in biological studies. Its inherent self-calibration in time domains provides quantitative insights through mechanisms such as Förster Resonance Energy Transfer (FRET). However, applying this technique in vivo introduces complexity: factors like attenuation and fluctuations in the biological environment can significantly affect the collected luminescence intensity from fluorophores. One emerging solution is to shift the emission wavelength to the near-infrared region, where reduced tissue scattering enables deeper penetration. Notably, lanthanide (III) emission centres offer tunable and stable luminescence in the 800 nm to 2000 nm range. Their typical microsecond-range lifetimes enable the exclusion of shorter-lived background signals (autofluorescence). These properties make lanthanide-based probes ideal candidates for in vivo time-resolved multiplexing and detection (Figure 1).

This report highlights my previous efforts to translate near-infrared-emitting lanthanide nanoprobes for deep-tissue, lifetime-based multiplexing and in vivo sensing. Specifically, I systematically explored strategies to optimize time-gated detection for multiplexing, improve probe efficiency<sup>[1]</sup>, develop bioconjugation methods that enable interference-free quantification<sup>[2]</sup>, and integrate these probes into advanced in vivo applications such as reactive oxygen species (ROS) detection<sup>[3]</sup> and temperature mapping<sup>[4]</sup>. These proof-of-concept studies lay a solid foundation for broader in vivo identification and quantification of other analytes, thus benefiting potential diagnostic and clinical translational applications.



Figure 1: Schematic illustration of time-resolved luminescence detection and imaging of lanthanide-based nanoprobes

## **References:**

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