

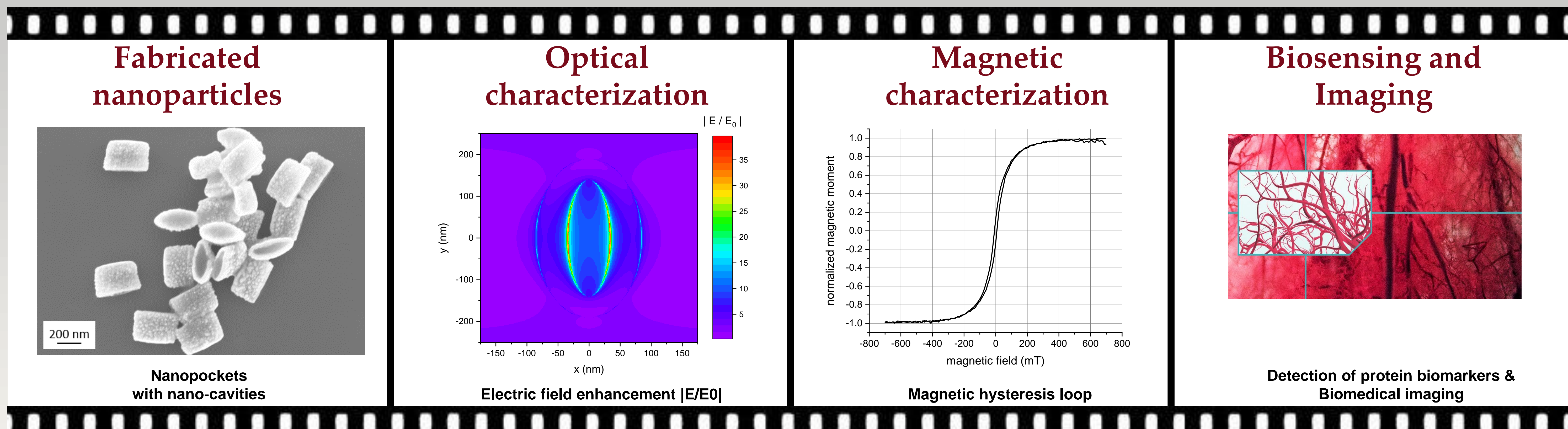
S. Schrittwieser^(a), M. Haslinger^(b), T. Mitteramkogler^(b), M. Mühlberger^(b), A. Shoshi^(c), H. Brückl^(c), M. Bauch^(d), T. Dimopoulos^(d), B. Schmid^(a), J. Schotter^(a), M. Eggeling^(a), C. Derntl^(a)

(a) AIT Austrian Institute of Technology, Molecular Diagnostics, Vienna, Austria. stefan.schrittwieser@ait.ac.at

(b) Profactor GmbH, Steyr, Austria

(c) Danube University Krems, Department for Integrated Sensor Systems, Wiener Neustadt, Austria

(d) AIT Austrian Institute of Technology, Energy Conversion & Hydrogen, Vienna, Austria



Introduction

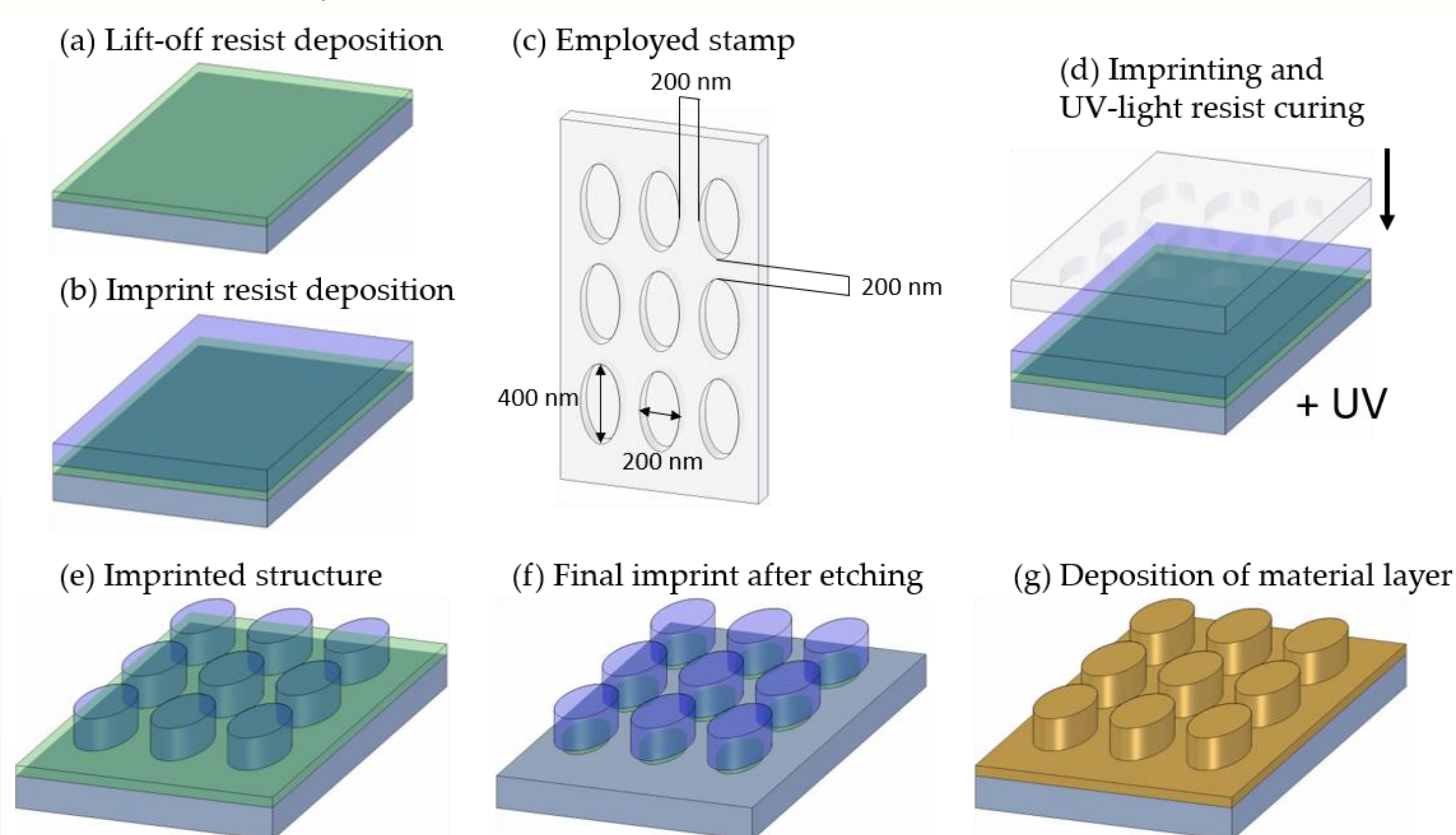
Nanostructured surfaces and nanoparticles are already widely employed in many different fields of research, and there is an ever-growing demand for novel types of nanoparticles. This is especially valid for multifunctional nanoparticles with physical properties that match the needs of a specific application. Additionally, there is an increasing need for improved medical imaging and fast and sensitive biosensing technologies.

We present the fabrication of three types of nanomaterials. Specifically, we present surfaces comprising a highly uniform array of elliptical pillars, nanoparticles with the shape of elliptical nanoplatelets as well as nanoparticles with the shape of nanopockets, possessing nano-cavities.[1,2] Optical and magnetic characterization reveals multifunctional properties of the fabricated nanomaterial. Furthermore, we report on the application of nanoparticles as probes for biosensing and as contrast agent for medical imaging.

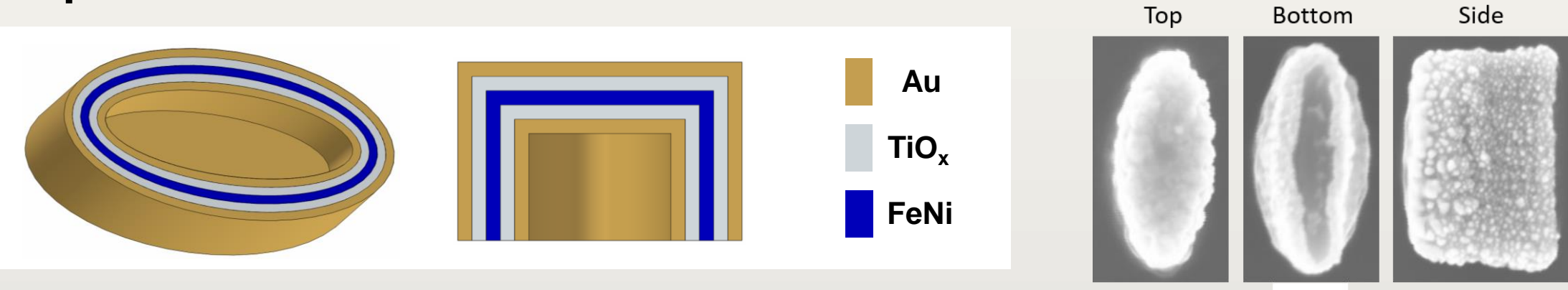
Nanoparticle fabrication

The structures are fabricated by nanoimprint lithography, physical and wet-chemical etching and sputter deposition of thin films of various materials to achieve a novel multifunctional nanomaterial with defined optical and magnetic properties.

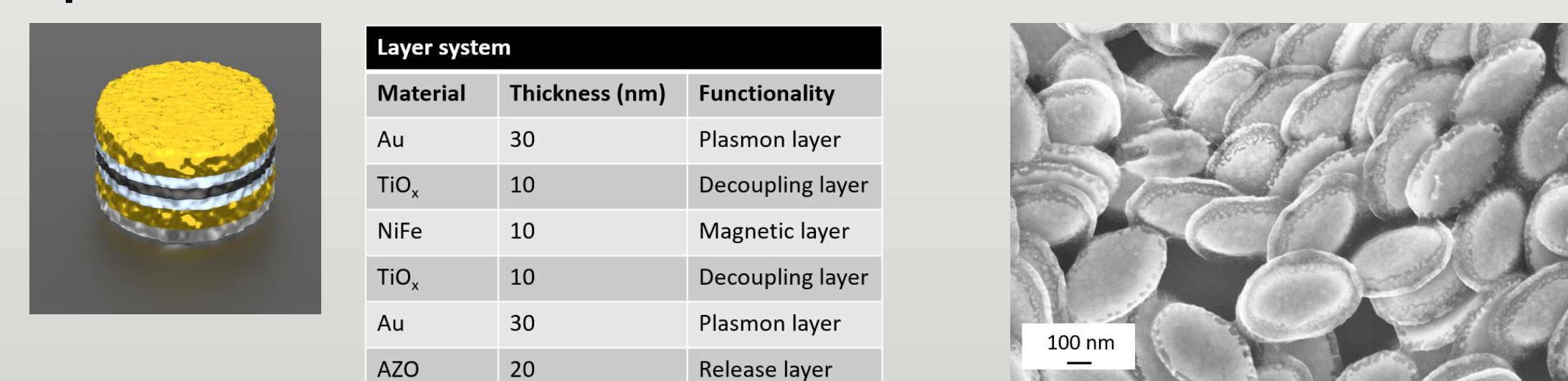
Scheme of fabrication process:



Nanopockets:

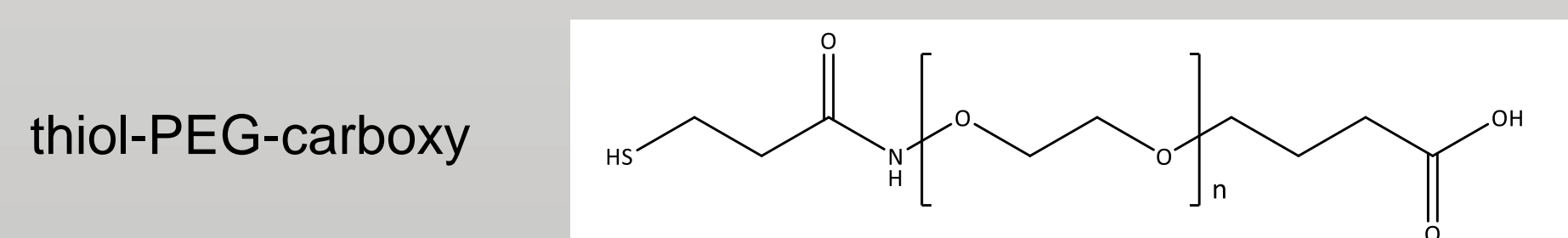


Nanoplatelets:

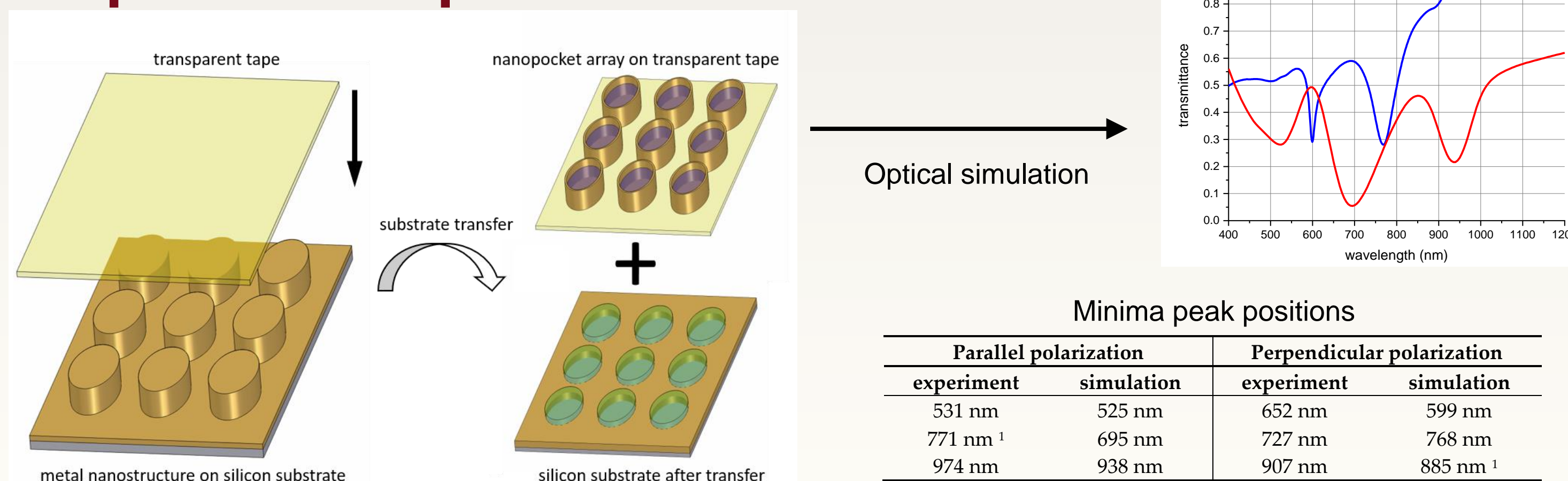


Surface modification and dispersion stabilization

Use of an additional linker molecule to stabilize aqueous nanoparticle dispersions and to allow for carbodiimide crosslinker chemistry for antibody binding (EDC/S-NHS).[1]



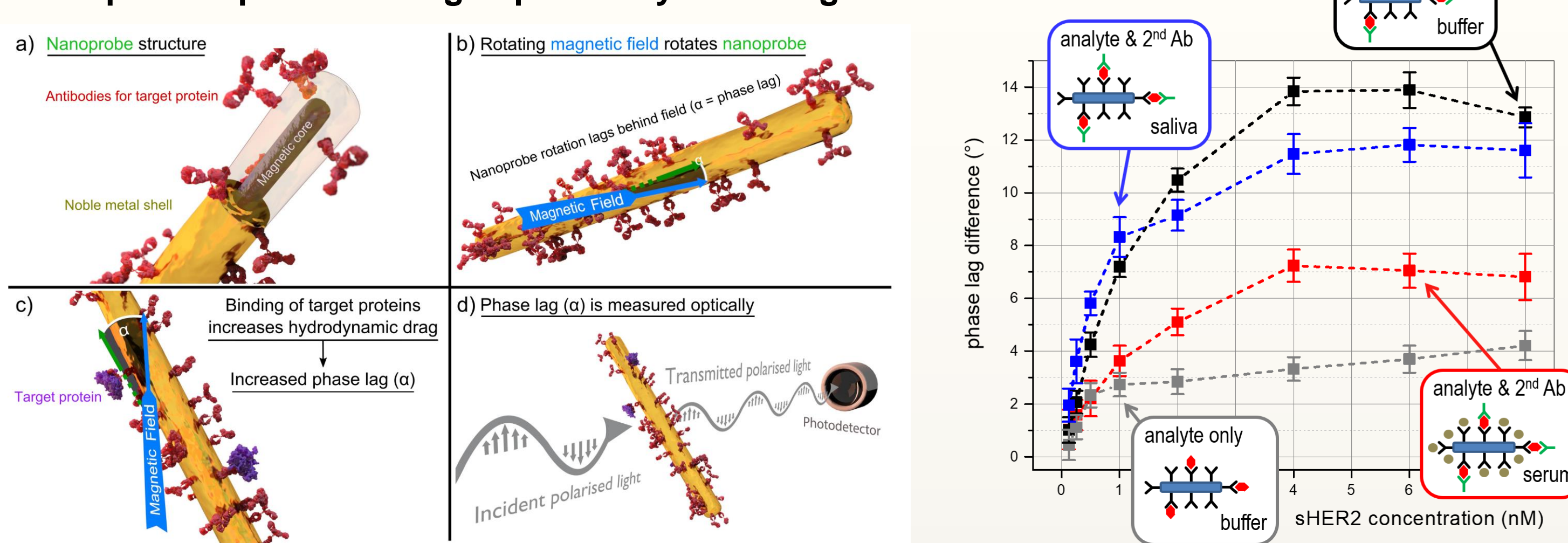
Optical nanoparticle characterization



Optical properties can be tailored to specific demands by predictive modeling steps

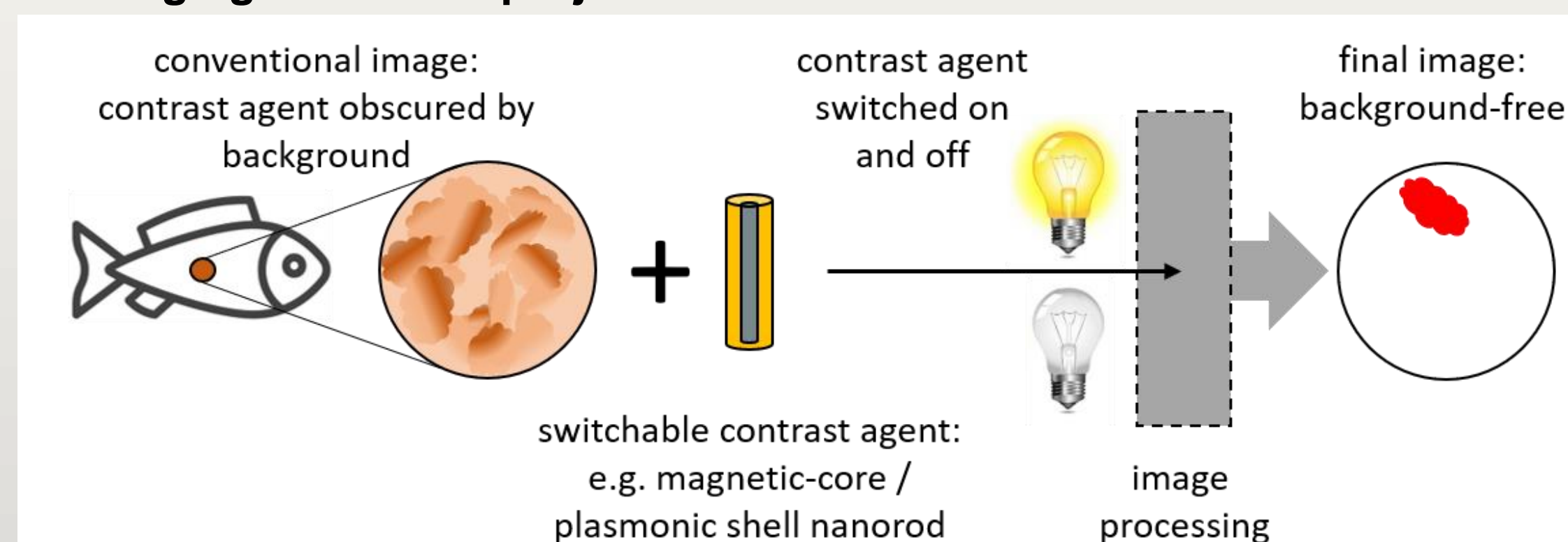
Biosensing and Imaging

Biosensing principle: Rotational dynamics of magnetic nanoparticle probes change upon analyte binding

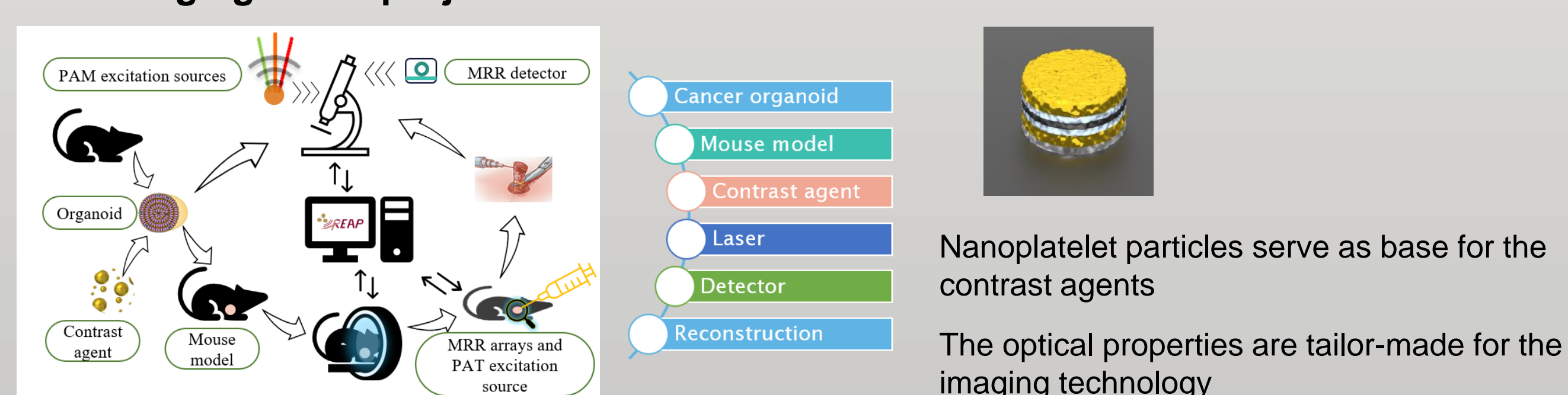


The breast cancer biomarker sHER2 (soluble domain of the transmembrane protein HER2) can be detected in complex sample solutions of saliva and serum if secondary antibodies (2nd Ab) are added to the assay. The extrapolated limit of detection is in the lower nM regime (e.g. 1.7 nM for saliva).[3,4,5]

Biomedical imaging SWIMMOT project



Biomedical imaging REAP project



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European Union's Horizon 2020 research and innovation programme under grant agreement number 899612 "SWIMMOT"; Photonics Public Private Partnership (PPP): H2020-ICT-2020-2 under grant agreement number 101016964 "REAP"; European Community's 7th Framework Programme under grant agreement number 246479 "NAMDIATREAM"; Austrian Research Promotion Agency (FFG) under grant agreement number 861414 "LAMPION".

<http://www.swimmot.researchproject.at>

<https://www.projectreap.eu/>

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- [2] S. Schrittwieser, M.J. Haslinger, T. Mitteramkogler, M. Mühlberger, A. Shoshi, H. Brückl, M. Bauch, T. Dimopoulos, B. Schmid, J. Schotter, *Nanomaterials* 9 (2019) 1790.
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