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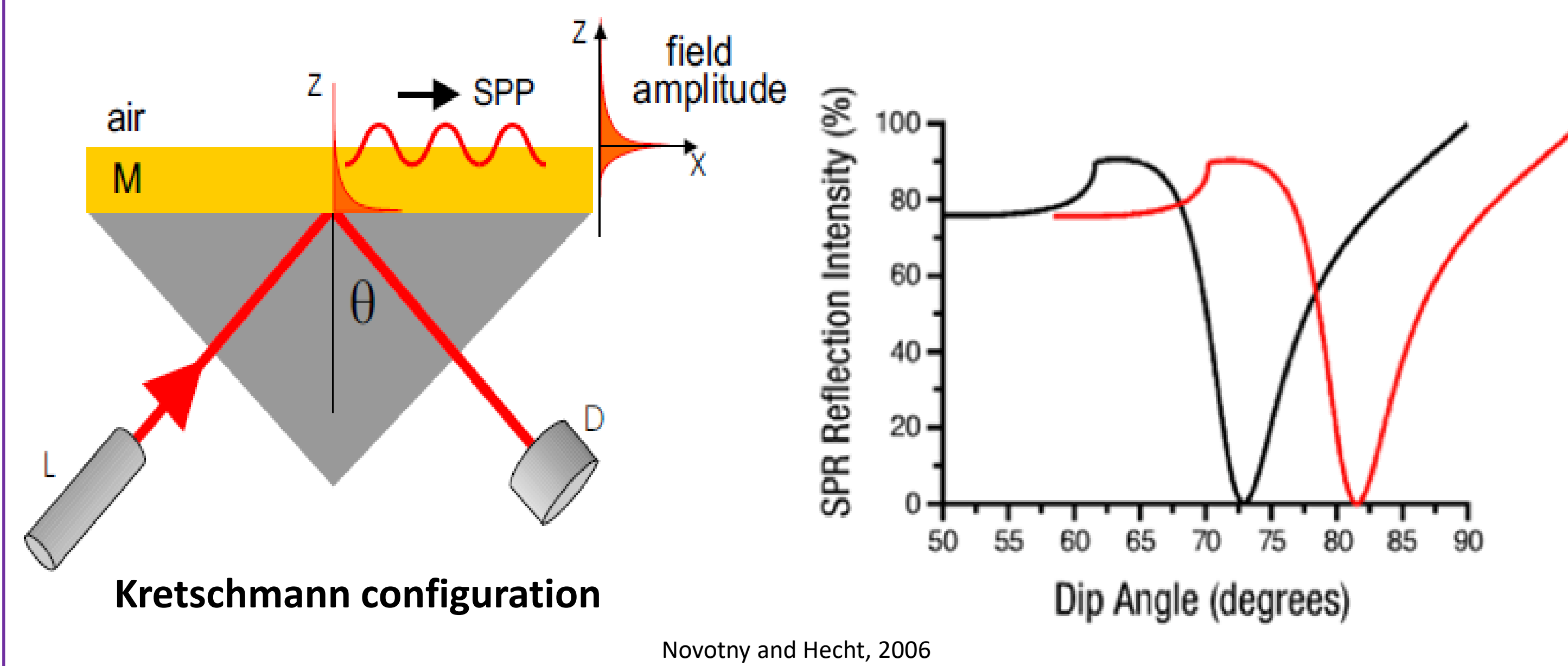


Background

Typical concentrations of physiological markers, such as cancer markers or cardiac markers can be very low (i.e. nano-molar range or below). Detecting a low concentration of biomolecules in the presence of an overwhelming background of natural constituents like blood components, becomes a very difficult task to achieve.

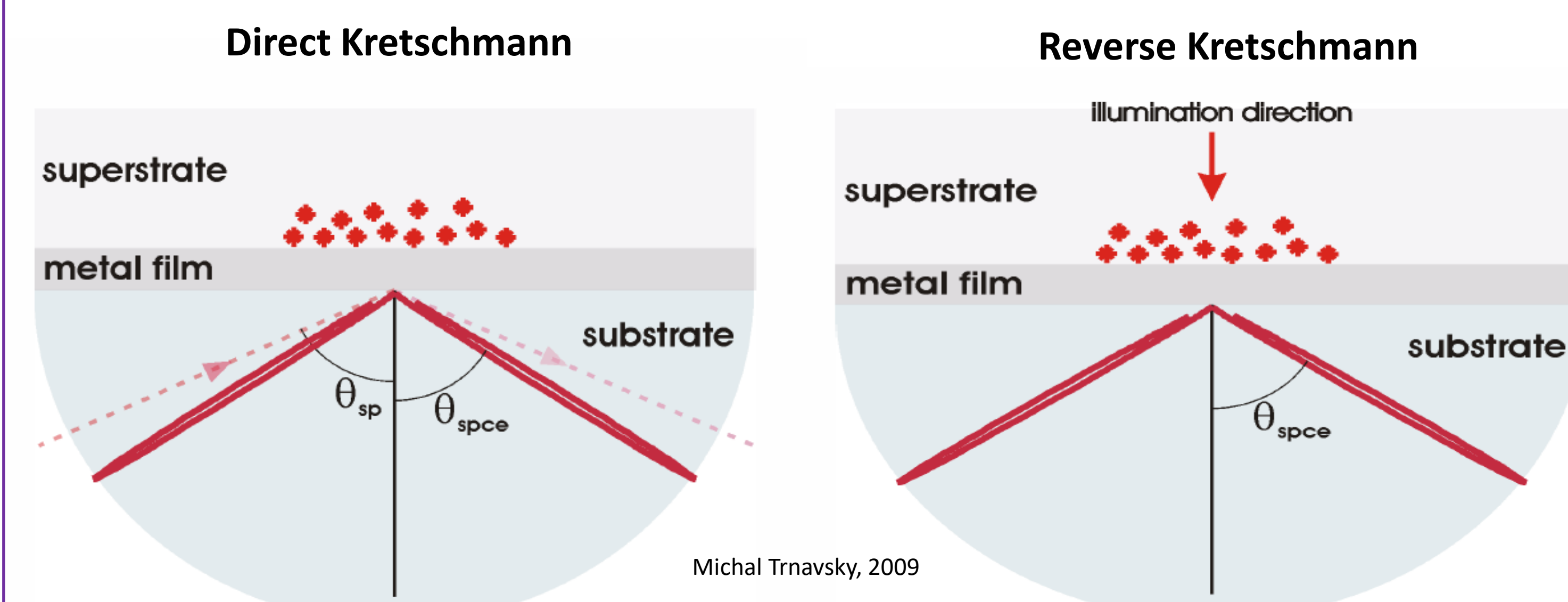
Fluorescence detection potentially offers the highest detection sensitivity. However, this is very much limited by **unwanted background signal** of naturally occurring physiological components. SPCE can be a powerful tool to detect low concentrations of biomolecules as well as molecular binding with **superior sensitivity**. The two crucial characteristics of SPCE are **surface confinement** for **fluorescence coupling** and **directional emission**.

Surface Plasmon Resonance: Surface charge density oscillation



SPR merely detects a change in refractive index. It does not distinguish non specific binding.

Surface Plasmon Coupled Emission: Opposite of SPR



Excitation via SPR

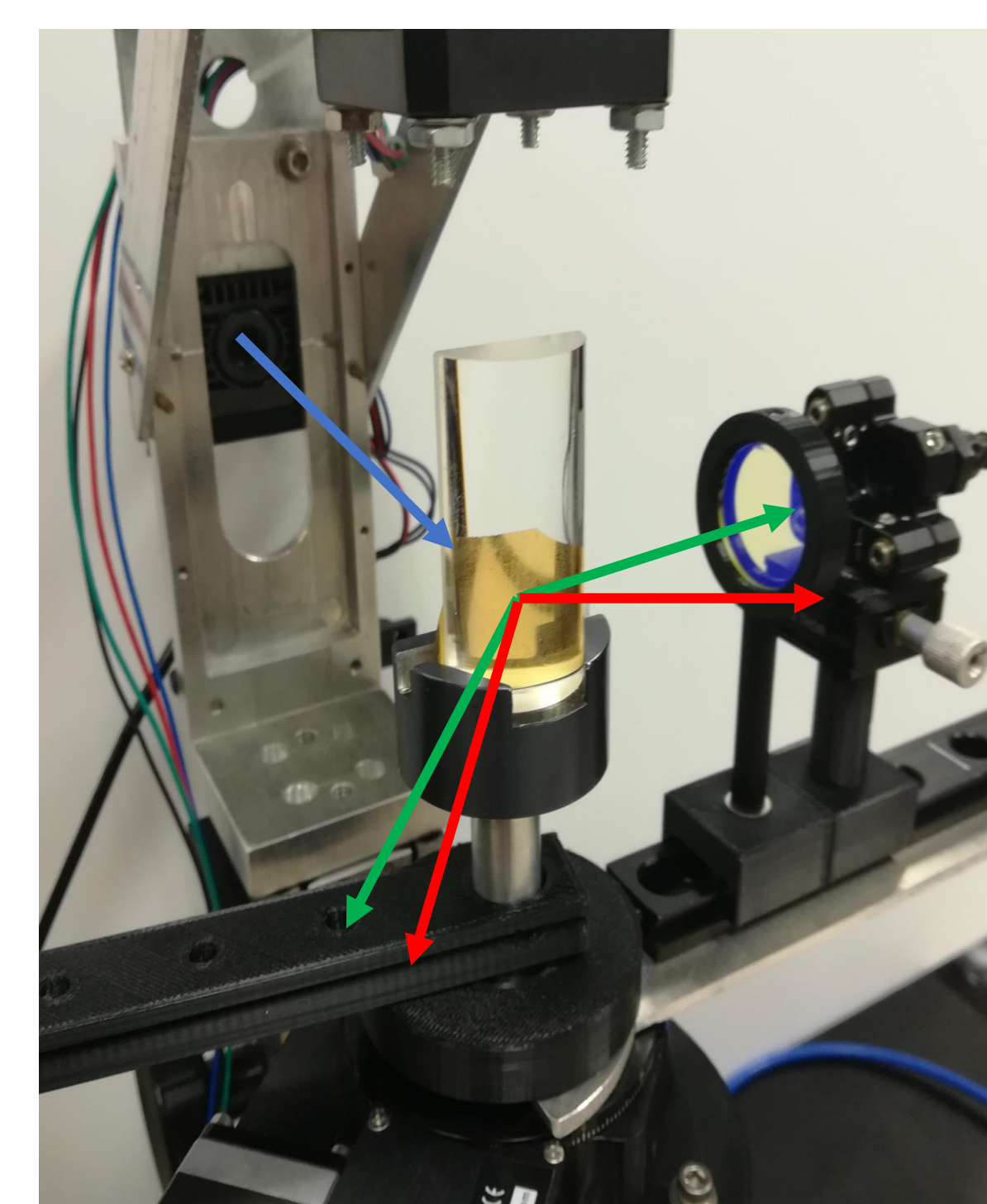
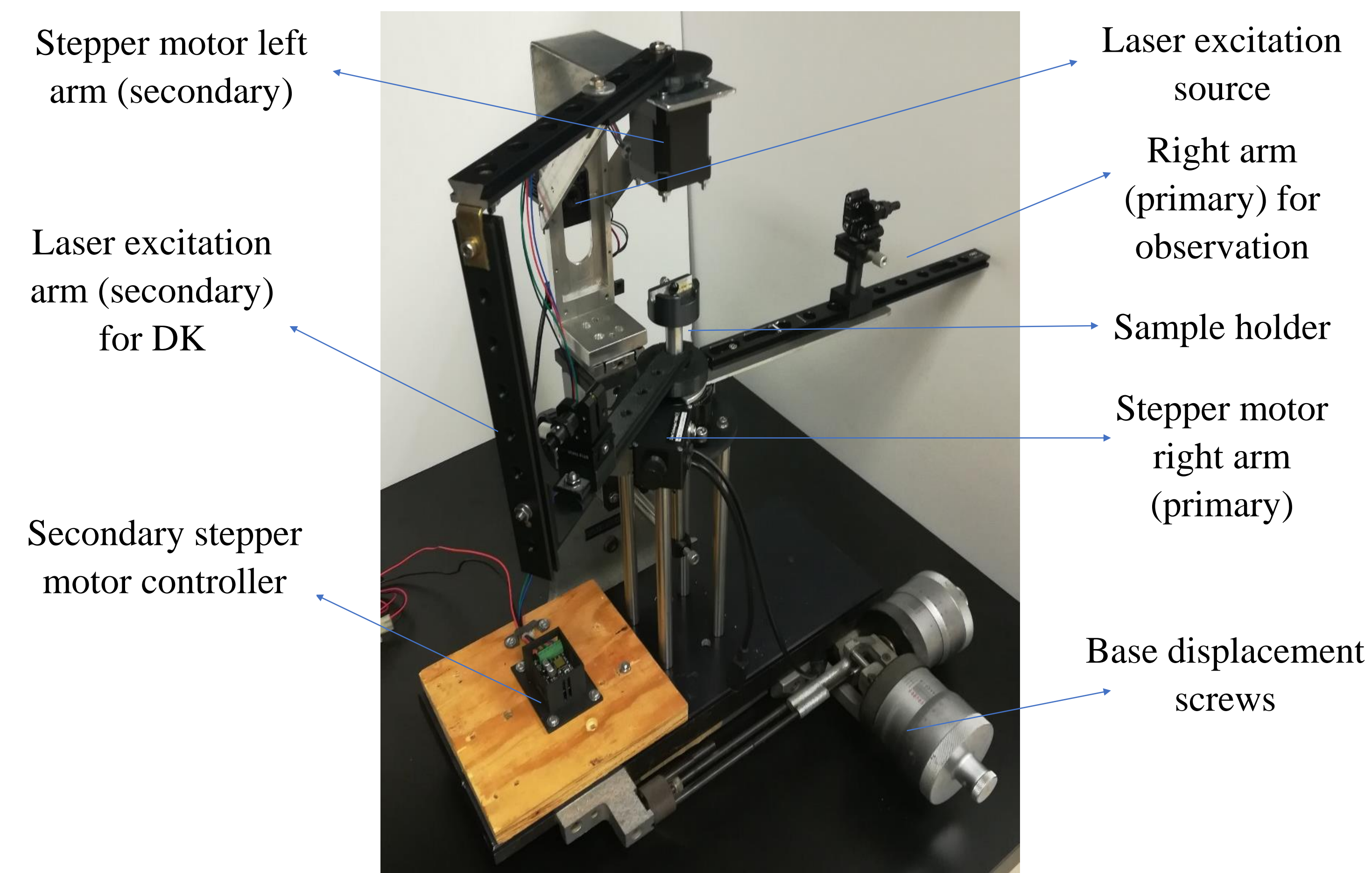
Excitation via external source

Smaller detection volume
More experimentally difficult

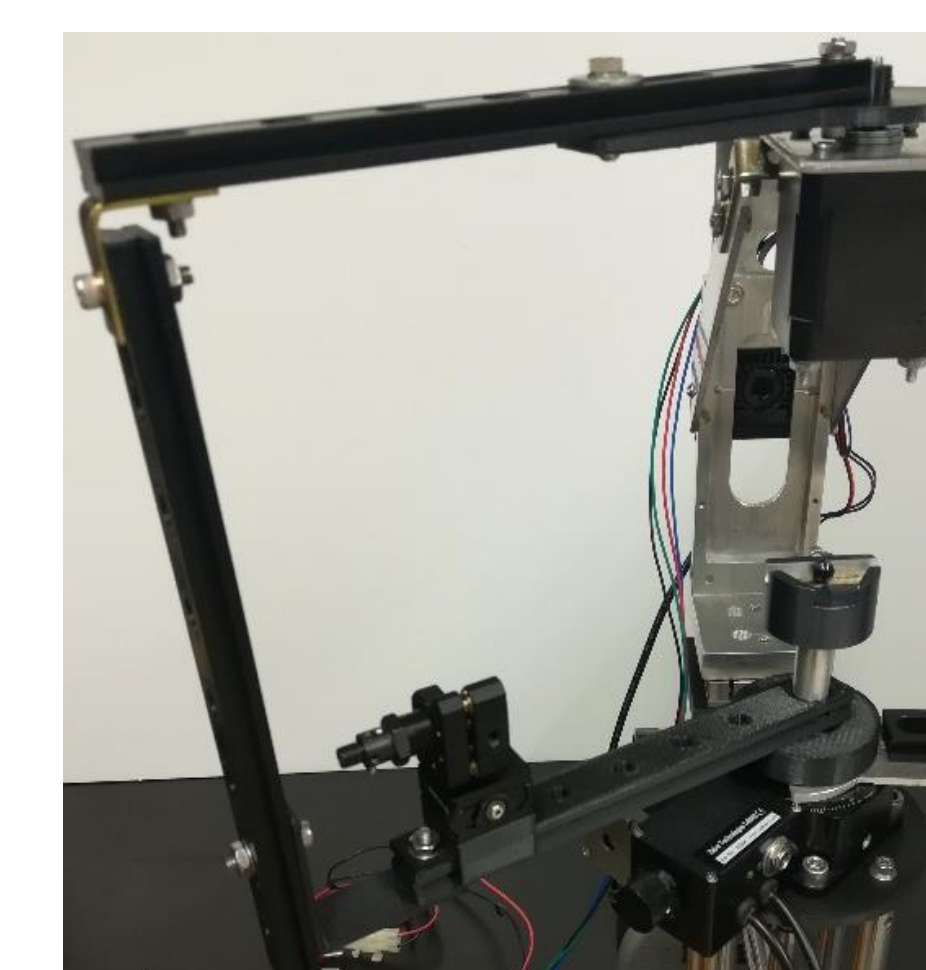
Experimentally easier
Not as "precise"

Experimental setup

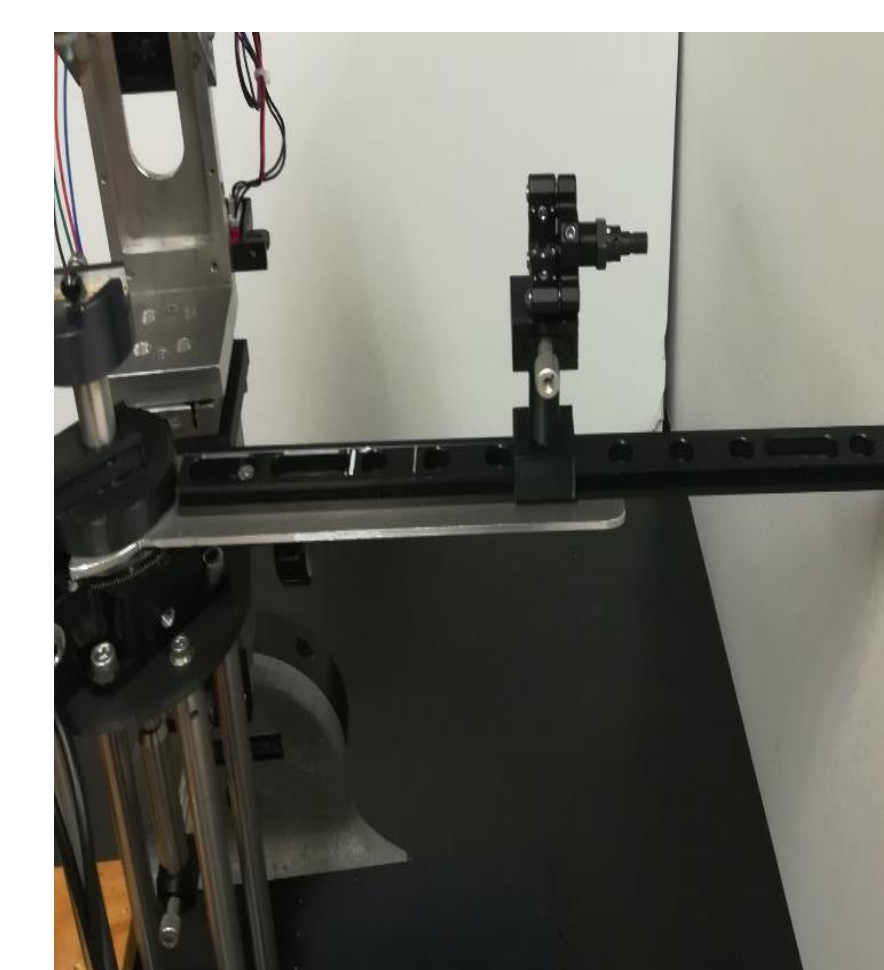
SPCE testing unit



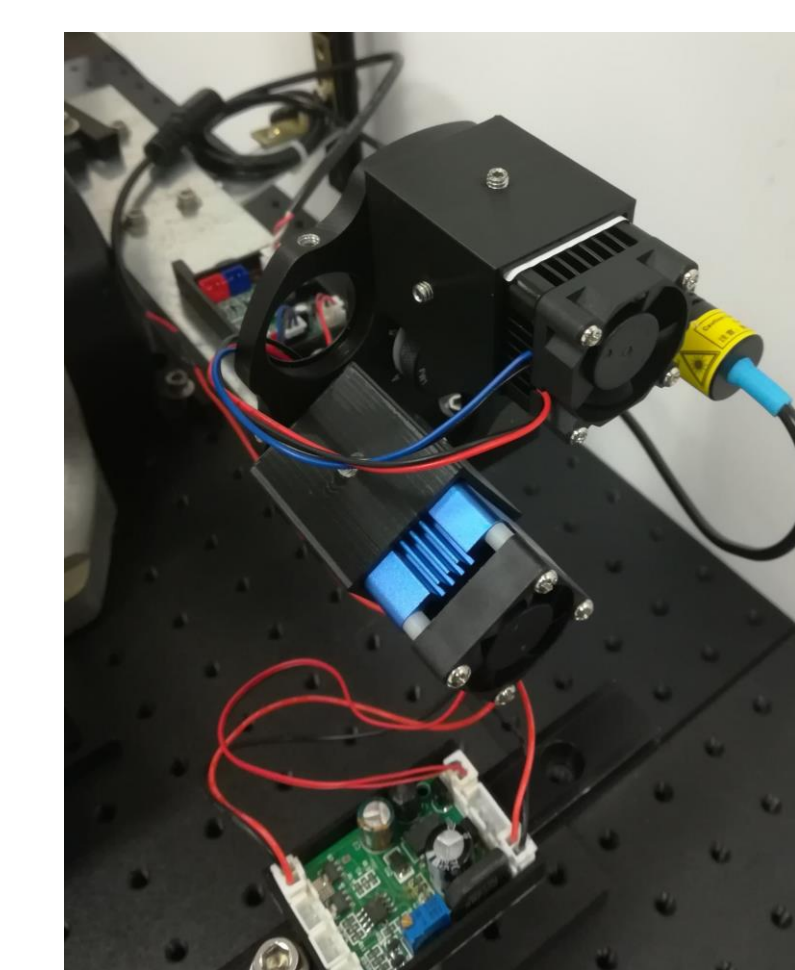
- Precise rotational motion around the sample
- Versatility for direct and reverse Kretschmann configuration
- Multi-laser excitation
- Easy control of sample position



Secondary arm (left)
Excitation arm for DK

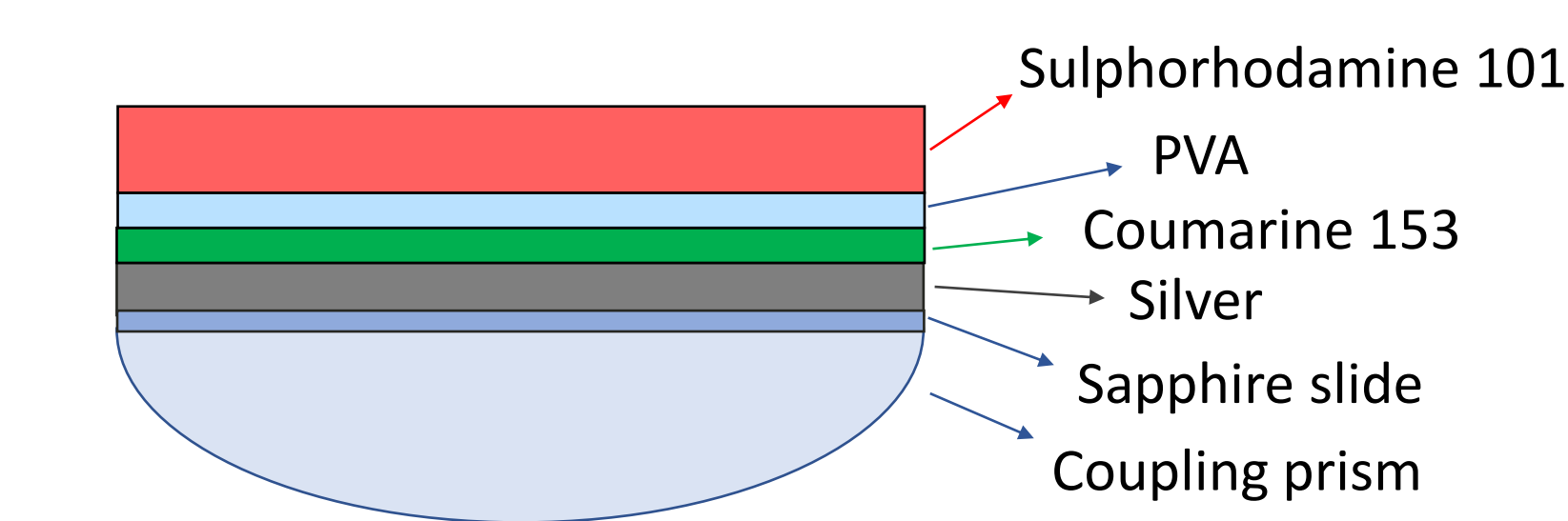
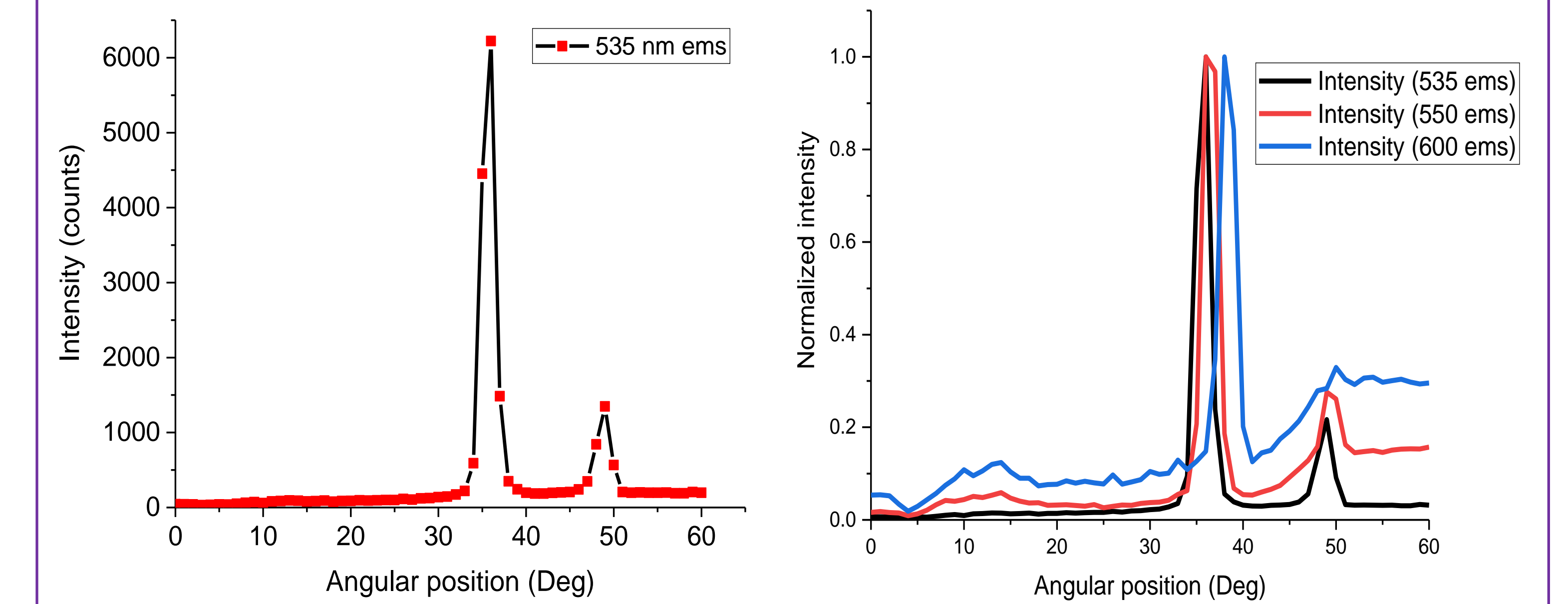


Primary arm (right)
Detection arm for RK and DK



Multi-laser excitation wheel

Results



- Test sensitivity and intensity of DK and RK
- Confirm role of distance dependent coupling

