# Three-dimensional Phase Imaging with Near Infrared Synchrotron Beam Using Phase-Retrieval Algorithm



Molong Han<sub>1</sub>,<sup>†</sup>, Daniel Smith<sub>1</sub>,<sup>†</sup>, Soon Hock Ng<sub>1</sub>, Tomas Katkus<sub>1</sub>, Aravind Simon John Francis Rajeswary<sub>2</sub>, Periyasamy Angamuthu Praveen<sub>2</sub>, Mark J Tobin<sub>3</sub>, Jitraporn Vongsvivut<sub>3</sub>, Saulius Juodkazis<sub>1,4</sub> and Vijayakumar Anand<sub>1,2</sub>,\*

10ptical Sciences Center, Swinburne University of Technology, Melbourne 3122, Australia; 2 Institute of Physics, University of Tartu, W. Ostwaldi 1, 50411 Tartu, Estonia; 3 Infrared Microspectroscopy (IRM) Beamline, ANSTO – Australian Synchrotron, Clayton, Victoria 3168, Australia;

4WRHProgramInternational Research Frontiers Initiative (IRFI) Tokyo Institute of Technology, Nagatsutacho,

Midoriku, Yokohama 226-8503, Kanagawa, Japan

<sup>+</sup>The authors contributed equally to this work; \*vijayakumar.anand@ut.ee;



## SUMMARY

- Near-infrared (NIR) synchrotron beam has been extracted and used for three-dimensional (3D) phase imaging.
- A pinhole was inserted in the path of the fork shaped NIR synchrotron beam and the Airy diffraction pattern was aligned with biochemical samples
- The diffracted intensity distribution was captured using an image sensor sensitive to NIR.
- A phase retrieval algorithm was used to estimate the 3D phase distribution from the recorded intensity distribution

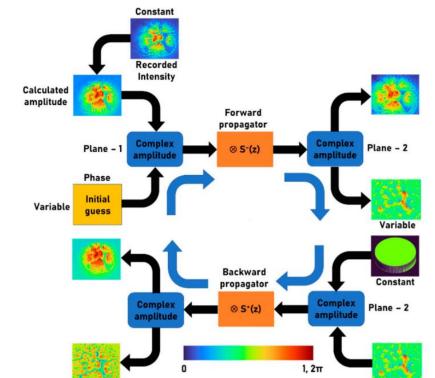
## EXPERIMENT

## Optical configuration of the IRM system

- IR beam extracted from the storage ring using a gold-coated mirror with a central slit has a fork shaped intensity distribution, which is focused using a Schwarzschild condenser on the sample[1]
- After passing through the sample, the beam is refocused by a Schwarzschild IR reflecting objective
- The beam then is collected by an MCT detector used for imaging in the single pixel scanning mode
- Visible light is also aligned collinearly with the synchrotron-IR beam for alignment of the sample during beam alignment in the microscope



- There are two planes of interest namely sample plane and sensor plane with two complex amplitudes  $\psi 1$  and  $\psi 2$  respectively with their intensities either known or measurable
- Initially, assuming phase variable is zero at detector plane, the complex amplitude ψ1 can be simply expressed as *VIo* (phase = 0). The complex amplitude at specimen plane ψ2 can be obtained by convolution of ψ1 and a forward spherical propagator S-(z) = exp[-j2πR/λ].
- The resulting complex amplitude's magnitude is replaced by the Airy intensity distribution or a constant valued matrix and the phase is retained.
- This modified complex amplitude is propagated to the sensor plane using a spherical propagator  $S+(z) = \exp[j2\pi R/\lambda]$
- The process will be repeated until estimation of phase images



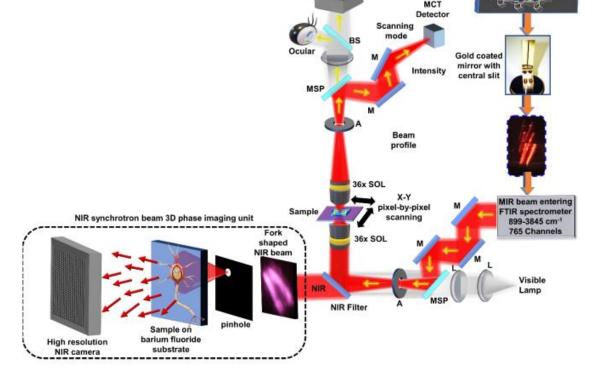


Figure 1: Schematic of the Australian synchrotron's IRM system and the attached synchrotron NIR phase imaging module (shown in the dotted line box). [4]

#### Attached NIR phase imaging module

- The high frequency filter is used to remove high-frequency components and extract the NIR beam.
- For getting a uniform intensity distribution, a pinhole was aligned with the maximum intensity region NIR beam and the Airy diffraction pattern was used to illuminate the biochemical sample.
- The phase information was extracted from this recorded intensity distribution using the phase-retrieval algorithm [5-7].



Figure 2 Schematic of the phase-retrieval algorithm[4]

#### RESULTS

The results obtained for two samples: randomly arranged latex beads with an average diameter of  $15\mu m$  and an insect wing

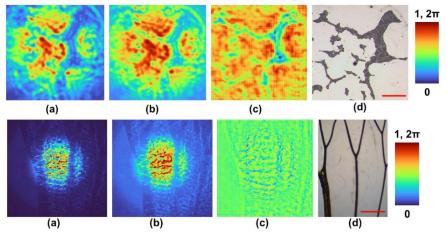


Figure 3: phase imaging results of both latex beads sample (top) and insect wing sample (bottom): (a) Intensity image, (b) amplitude image, (c) phase image, and (d) reference image of the bead samples. [4]

### CONCLUSION

- A single shot, interferenceless phase imaging technique has been developed and demonstrated in the IRM system of the Australian Synchrotron for the first time.
- The phase-retrieval used in this approach has a rapid convergence of 2-4 iterations which may allow real-time phase imaging with only a slight decrease in the temporal resolution

#### **References:**

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