

PROCEEDINGS OF SPIE

[SPIDigitalLibrary.org/conference-proceedings-of-spie](https://spiedigitallibrary.org/conference-proceedings-of-spie)

Ultrafast laser scanning cellular microscopy by spatiotemporally encoded virtual sources

Wenwei Yan
Jianglai Wu
Kenneth K. Y. Wong
Kevin K. Tsia

SPIE.

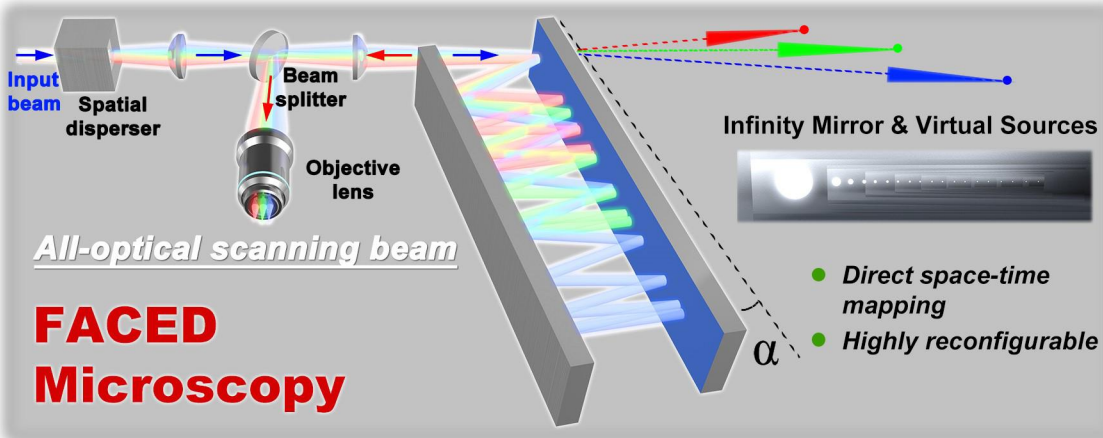
Ultrafast laser scanning cellular microscopy by spatiotemporally encoded virtual sources

Wenwei Yan, Jianglai Wu, Kenneth K.Y. Wong and Kevin K. Tsia

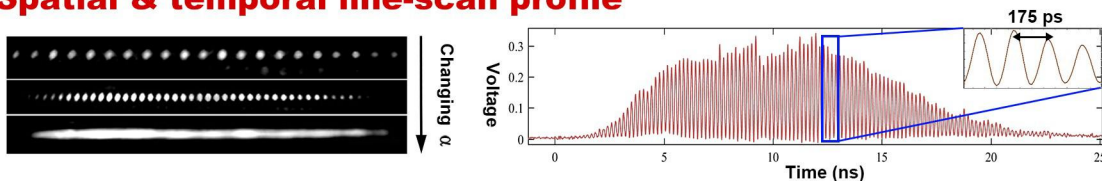
Department of Electrical and Electronic Engineering, The University of Hong Kong



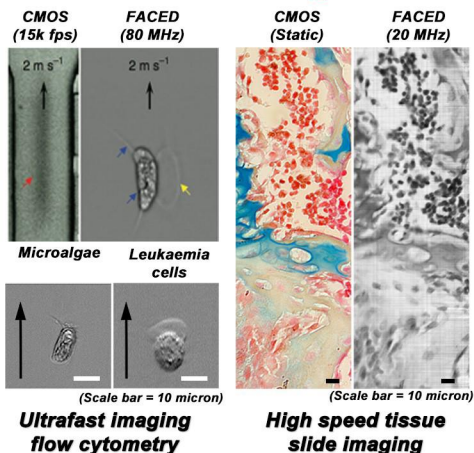
Introduction We report a new type of **all-optical ultrafast laser-scanning microscopy** (at a line-scan rate of 20 MHz) based on a phenomenon called *free-space angular-chirp-enhanced delay* (**FACED**). It results in the generation of a reconfigurable array of spatiotemporally encoded virtual pulsed sources, which acts as a scanning laser beam. We demonstrate its application in high-throughput multivariate image-based single-cell analysis (10,000 cells/sec).



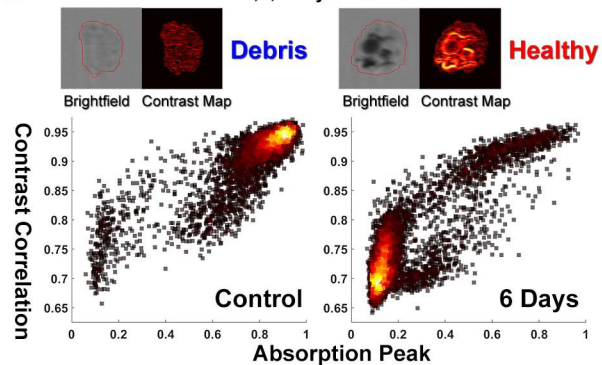
Spatial & temporal line-scan profile



From ultrafast single-cell imaging to analysis



- High speed microfluidic flow (>1 m/s)
- Human leukaemia cells labelled to visualize lysosome
- Nutrient starvation for 2,4,6 days



References:

[1] Wu, J., et al., "Ultrafast laser-scanning time-stretch imaging at visible wavelengths," *Light: Science & Applications* 6, e16196 (2017).