

## Integrated photonic pillar scatterers for speeding up classification of cell holograms

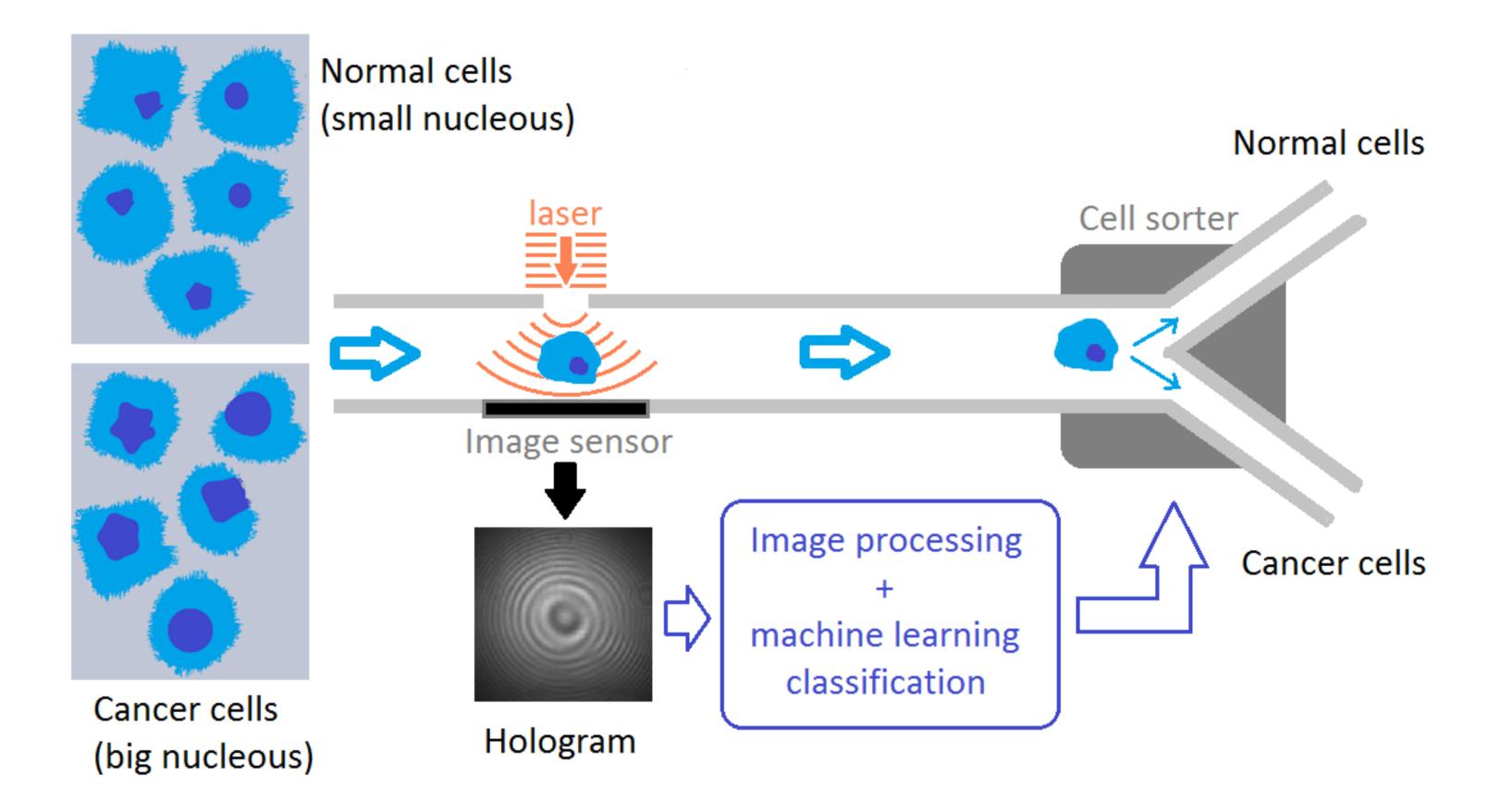
Alessio Lugnan<sup>1</sup>, Joni Dambre<sup>2</sup>, Peter Bienstman<sup>1</sup> <sup>1</sup>Photonics Research Group, UGent – imec, Technologiepark 15, 9052 Gent, Belgium <sup>2</sup>IDLab, UGent - imec, Technologiepark 15, 9052 Gent, Belgium *Tel:* +32 9 264 3450, *Fax:* +32 9 264 3593, *e-mail:* alessio.lugnan@ugent.be

**Flow cytometry** enables *high-speed sorting* of different kinds of cells flowing in a fluidic channel

employing

Digital holographic microscopy

- Obtains information about the cell optical structure by lighting it with coherent light and acquiring its interference pattern (*hologram*)
- Allows for **label-free classification** without altering the cells, but **image reconstruction is computationally expensive** (major limit to sorting speed)



We designed a *passive integrated photonic stage* to speed up machine learning classification of cell holograms.

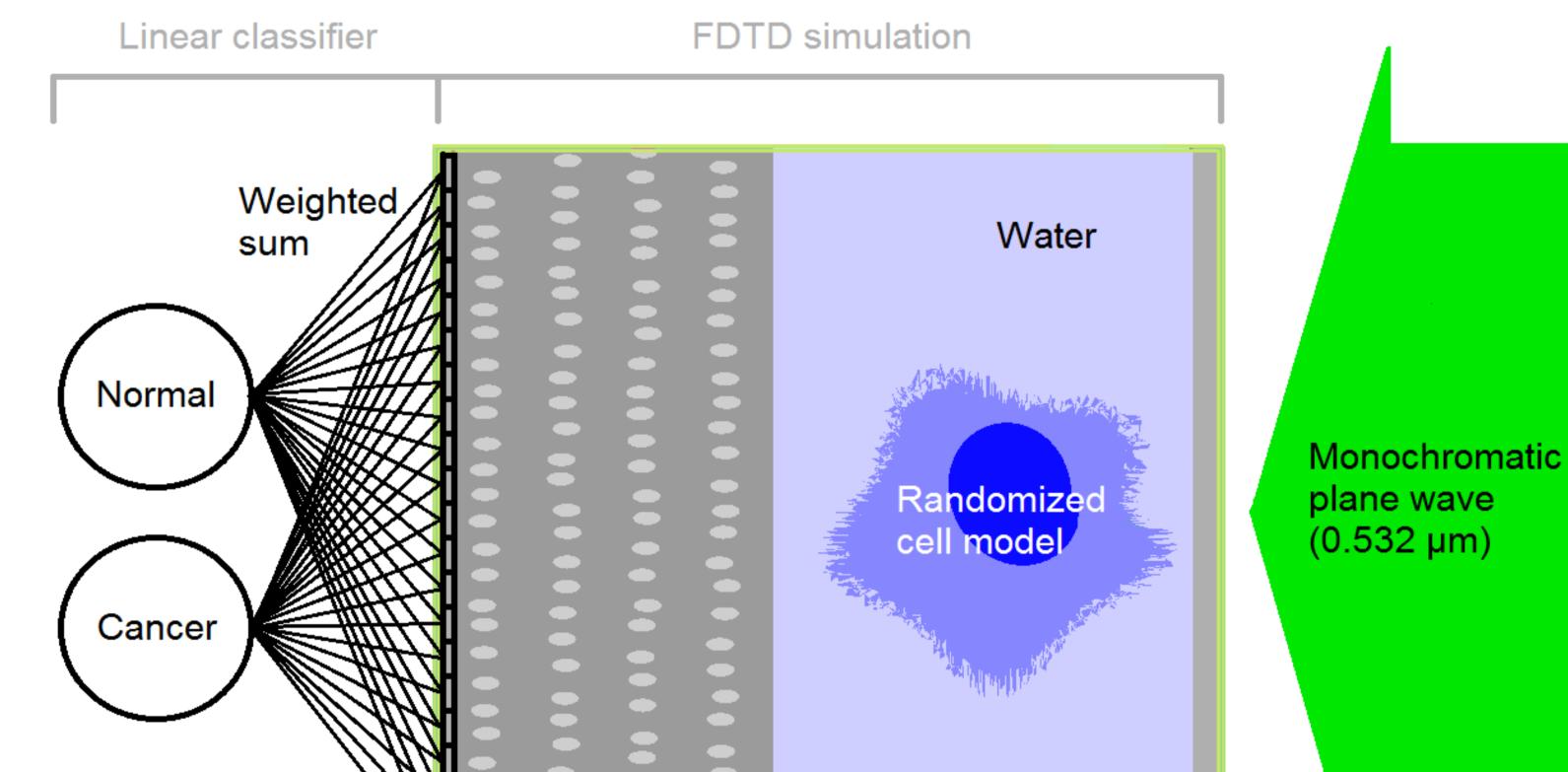
A cell hologram is determined by

- Small refractive index contrast: n(cell)~1.37 n(water)~1.34
- Negligible absorption

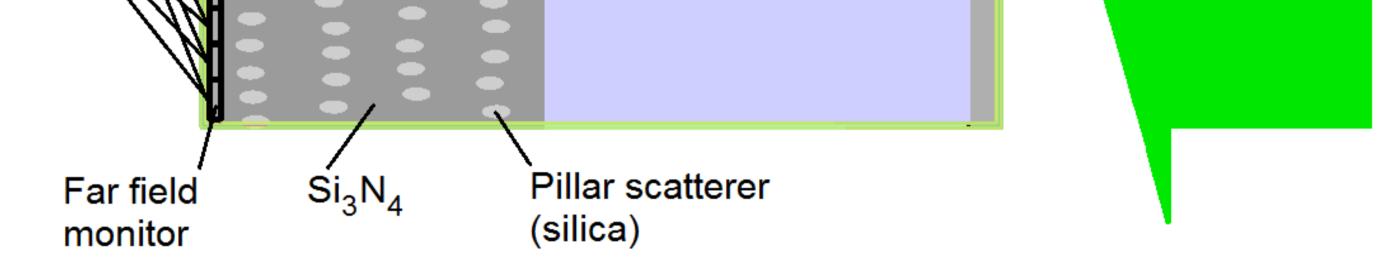
The information is encoded in the *optical phase* of the light scattered by a cell

The transfer function between the optical phase and the optical intensity, which is measured by a detector, is of a sinusoidal nature

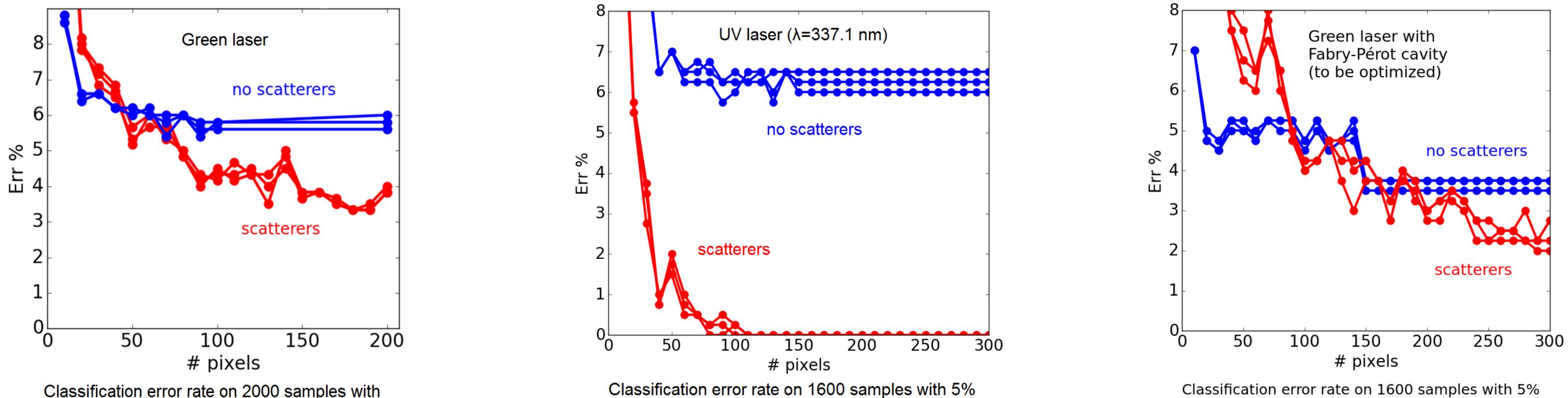
Power-independent nonlinearity available for computation

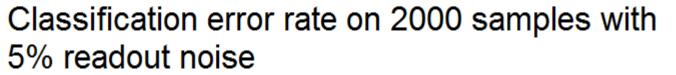


We employed a **spatial analog of reservoir computing**, in which the reservoir is a *collection of silica scatterers* that mixes the phase-encoded optical signal before applying a linear classifier.



- Simulations show that the application of scatterers increases the performances of a logistic regression in the classification of cells with two different average nucleus sizes ("normal" and "cancer" cells).
- In order to increase the phase-to-intensity nonlinearity with respect to different nucleus sizes, and thus the performances, the light wavelength can be decreased (UV laser) or the  $\bullet$ cell can be placed in an optical cavity (e.g. integrated Fabry-Pérot cavity using Bragg reflectors).





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Photonics Research Group, INTEC-Department, Ghent University-IMEC Center for Nano- and Biophotonics (NB-Photonics), Ghent University http://photonics.intec.ugent.be

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