

BISET: Biological Image Super-resolution Enhanced with Tensor

- ▶ MORPHEME team - I3S Lab, INRIA SAM - (**Eric Debreuve**, Laure Blanc-Féraud)
Image processing
- ▶ SIGNET team - I3S Lab, (Gérad Favier, **Henrique Goulard**)
Signal processing, Tensor
- ▶ iBV Lab, (Sébastien Schaub, Guillaume Sandoz)
Optics, biology (ion channels)

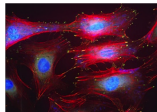
Academy 1 of UCA, 4, dec. 2017



institut Valrose
B i o l o g i e

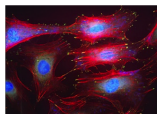
Fluorescence microscopy

- ▶ Structure of interest in living cells can be imaged through the microscope using fluorescent proteins
- ▶ Gene of fluorescent molecule can be combined with gene of proteins of structure we want to study
Nobel Prize of chemistry 2008
- ▶ Illumination with a laser causes the fluorophores to emit photons



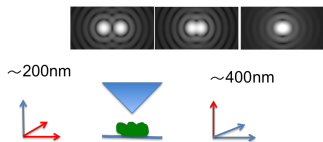
Fluorescence microscopy

- ▶ Structure of interest in living cells can be imaged through the microscope using fluorescent proteins
- ▶ Gene of fluorescent molecule can be combined with gene of proteins of structure we want to study
Nobel Prize of chemistry 2008
- ▶ Illumination with a laser causes the fluorophores to emit photons



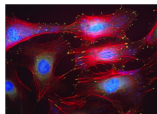
Resolution limits

- ▶ physical diffraction limit of optical systems
- ▶ 200nm lateral, 400 axial.



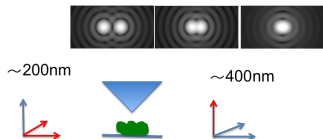
Fluorescence microscopy

- ▶ Structure of interest in living cells can be imaged through the microscope using fluorescent proteins
- ▶ Gene of fluorescent molecule can be combined with gene of proteins of structure we want to study
Nobel Prize of chemistry 2008
- ▶ Illumination with a laser causes the fluorophores to emit photons



Resolution limits

- ▶ physical diffraction limit of optical systems
- ▶ 200nm lateral, 400 axial.



- ▶ Study of ion channels, membrane proteins that allow the passage of ions through the cell membrane. Resolution needed less than 50 nm.

Multi-Angle TIRF \rightarrow axial super-resolution

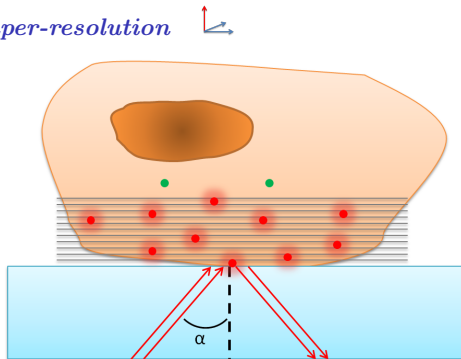
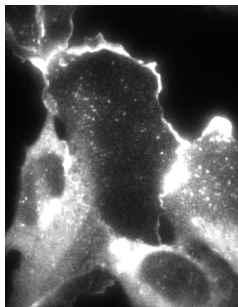


Figure: TIRF Acquisitions.

- ▶ TIRF *Total Internal Reflexion Fluorescence* Microscope is a commercial system
- ▶ **Multiple Angle** TIRF prototype at iBV, no more than 3 prototypes in France,
- ▶ Reconstruction algorithm for the depth information, **30nm of resolution over 400nm thick layer** over the coverslip.
- ▶ Method of choice to visualize processes around the plasma membrane

Figure: MA-TIRF Acquisitions.

- ▶ TIRF *Total Internal Reflexion Fluorescence* Microscope is a commercial system
- ▶ **Multiple Angle** TIRF prototype at iBV, no more than 3 prototypes in France,
- ▶ Reconstruction algorithm for the depth information, **30nm of resolution over 400nm thick layer** over the coverslip.
- ▶ Method of choice to visualize processes around the plasma membrane

3D super-resolution by mixing MA-TIRF and SIF

Optical axial Fluctuation \rightarrow *lateral super-resolution*



- ▶ use axial SIF *Super-resolution Intensity Fluctuation* (blinking):
Acquisition of 100 images in 1 second,
- ▶ each image is blurred and low level SNR (signal on noise ratio), but the temporal fluctuation of fluorescent molecule will be used to localize precisely the fluorophore
- ▶ analysis of the temporal image series: cumulant, source separation

Mixing MA-TIRF and SIF \rightarrow *Tensor analysis*

- ▶ MA-TIRF + SIF gives both axial and lateral super-resolution but big volume of 3D + t data to process
- ▶ Two possible tensor-based approaches:
 1. Separation of independent sources based on decomposing **high-order cumulants** of temporal image series
 2. Direct decomposition of MA-TIRF + SIF data: **joint exploitation of several diversities** for deterministic (blind) source separation.

3D super-resolution by mixing MA-TIRF and SIF

Optical axial Fluctuation \rightarrow *lateral super-resolution*



- ▶ use axial SIF *Super-resolution Intensity Fluctuation* (blinking):
Acquisition of 100 images in 1 second,
- ▶ each image is blurred and low level SNR (signal on noise ratio), but the temporal fluctuation of fluorescent molecule will be used to localize precisely the fluorophore
- ▶ analysis of the temporal image series: cumulant, source separation

Mixing MA-TIRF and SIF \rightarrow *Tensor analysis*

- ▶ MA-TIRF + SIF gives both axial and lateral super-resolution but big volume of 3D + t data to process
- ▶ Two possible tensor-based approaches:
 1. Separation of independent sources based on decomposing **high-order cumulants** of temporal image series
 2. Direct decomposition of MA-TIRF + SIF data: **joint exploitation of several diversities** for deterministic (blind) source separation.