abbelight now, we see

The future of imaging, from microscopy to single-molecule localization

abbelight Now, we see.





Preparation

Optimal sample preparation requires multi-disciplinary know-how



Imaging

High-quality and reliable results depend on precise instruments



Analysis

Handling of data requires powerful and tailored analytical tools

Single Molecule Localization Microscopy (SMLM) combines quantitative information with the highest resolution achievable in light microscopy, and is therefore a game changer in many biological studies.

To efficiently accommodate SMLM in a research workflow, it is crucial to build a diverse expertise ranging from sample preparation to image acquisition and data analysis.

abbelight offers are designed to guide and provide support with the best instruments, software and expertise, to speed-up the imaging workflow within your research applications.



abbelight research

Advice, samples and reagents for your projects



abbelight instruments

The best performances



abbelight data

An all-in-one software designed for acquisition, processing and analysis

abbelight Our solutions



abbelight research

Biological samples Reagents Advice, protocols and training



abbelight instruments

Evolutive add-ons for inverted microscopes

SAFe 180 - Large field of view TIRF and easy single-molecule imaging

SAFe 360 - High precision, multicolor 3D single-molecule imaging



abbelight data

Acquisition and real-time image reconstruction Advanced processing Easy visualization and analysis



abbelight About us

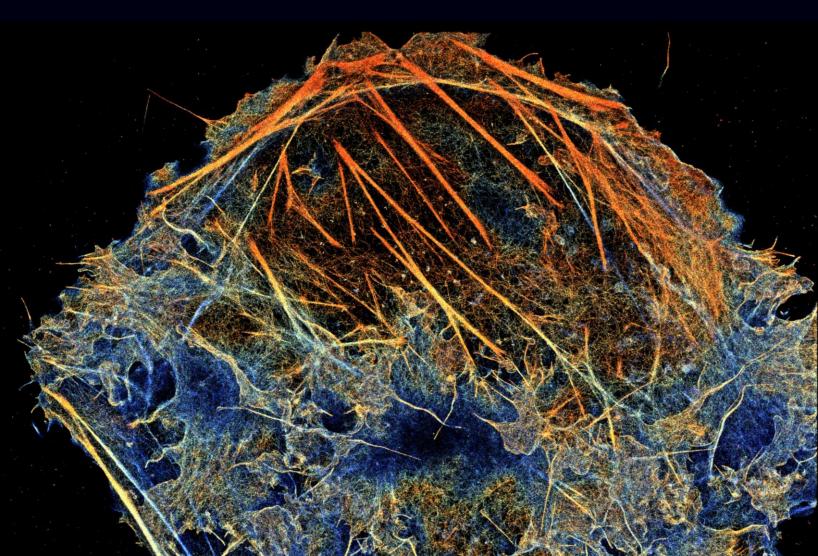
abbelight is the result of 10 years of academic research on cutting-edge detection methods in fluorescence microscopy; our team holds a vast pool of knowledge in every aspect of localization microscopy.

Still fully immersed in the research environment, abbelight converts the most advanced technology into robust, evolutive, and accessible products; its software developers make them fast, efficient and intuitive to use for their clients.

Our focus

Develop features that matter for research and researchers:

- Speed up not only data acquisition, but the full imaging workflow
- Ensure transparency about product and technology
- Produce reliable and reproducible data with fewer artifacts not just better images
- Reveal and fully exploit data complexity





SAFe accelerator by researchers for researchers

Our team of scientists is involved in multi-disciplinary research, education and international training.

With our SAFe accelerator we provide time, knowledge and expertise to fast-track the imaging workflow and overcome experimental hurdles during the first steps of a new project.



SAFe samples and reagents for calibration and training

Cell type

- Cos7
- U2OS
- HeLa
- HEK
- primary neurons
- macrophages
- Escherichia coli

Cell structure

- microtubules and actin cytoskeleton
- spectrin cytoskeleton
- clathrin coated pits
- mitochondria
- podosomes
- bacterial cell wall and membrane

Fluorophores

- Atto 488, WGA-AF® 488
- AF [®] 532, CF [®] 532, Cy3b
- AF ® 555, AF ® 594, CF ® 555, AF ® 568, CF ® 568, Cy5, MemBrite™ 568, TMR
- AF [®] 647, CF [®] 647, AF [®] 680, CF [®] 680, MemBrite[™] 640, Actin-stain 670

Ready-to-use solutions for localization microscopy (dSTORM), compatible with the dyes listed above and more.

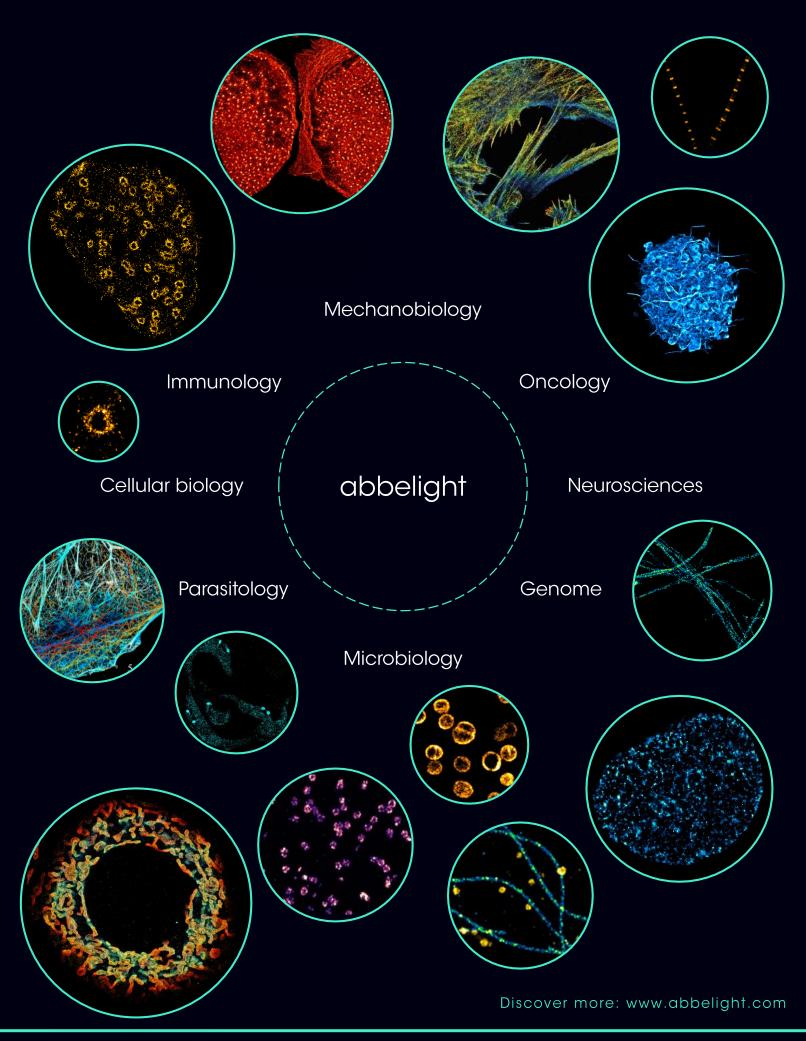
abbelight provides two solutions to induce photoswitching of fluorophores:

- SmartKit is a mounting medium for fixed cells
- Blinking Pad maintains non-adherent biological objects immobilized on the coverslip



- 10 doses per box
- 200 µL per dose
- 30 sec preparation
- 2 months in a fridge
- High efficiency and stability over time





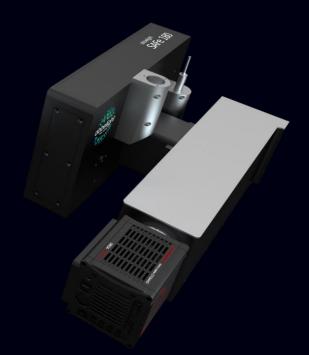




SAFe 180 for 2D multicolor single-molecule imaging with **10 nm** localization precision over the largest field of view of the market with **abbelight SAFe Light** technology.

- Super-resolution imaging modes: PALM, STORM, Single-Particle Tracking...
- Illumination modes: Epi, HiLo, TIRF
- 10 nm x 10 nm localization precision
- The largest homogeneous 2D field of view 150 x 150 µm²
- Low power laser needed thanks to abbelight SAFe Light technology
- Automatic switching between different illumination modes
- Fully supported by abbelight Data and abbelight Research





Adaptable

SAFe 180 and 360 can be adapted to any inverted microscope.

In a few minutes, you can add unique features to your existing microscopes.

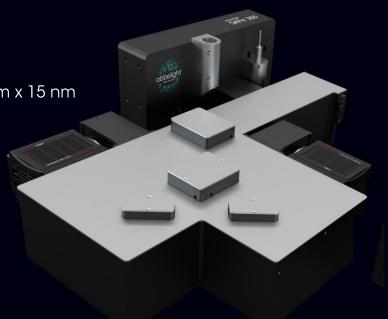
Add	TIRF	to your	Confocal
	PALM		Spinning-disk
	STORM		Widefield
	SPT		SIM
	PAINT		STED





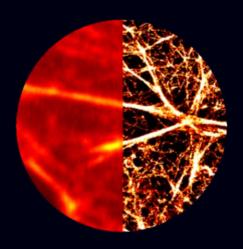
SAFe 360 to obtain 3D single-molecule information with **15 nm** localization precision in **z**, while maintaining the 10 nm localization precision in x and y.

- All **SAFe 180** features
- Ultimate 3D localization precision: 10 nm x 10 nm x 15 nm
- 1 µm capture range
- 5-10 µm imaging depth
- Absolute measurement relative to the coverslip for reproducible, artefact-free measurement
- Simultaneous multicolor imaging in Epi, TIRF, or single-molecule mode
- New extra modalities for live cell imaging

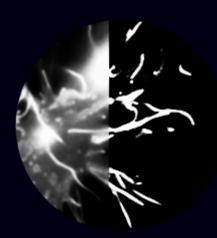


Multimodal

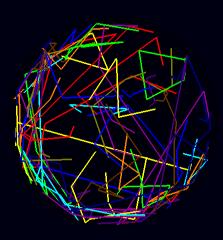
Fluorescence nanoscopy



Widefield / TIRF



Live imaging







NEO is abbelight's intuitive and user-friendly software. It allows the production and manipulation of 3D quantitative data. NEO software offers a complete workflow for all imaging modalities in Single Molecule Localization Microscopy: from acquisition to analysis.

Acquisition

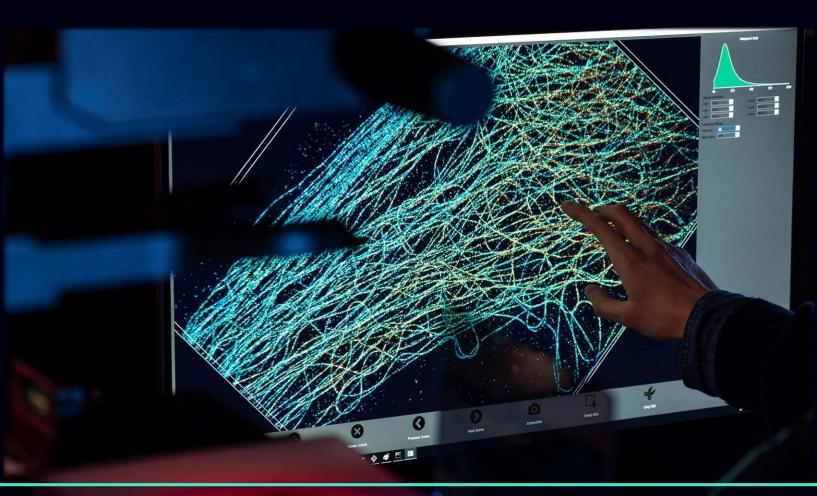
- Full instrument control
- Simultaneous multicolor imaging
- Acquisition guided by decision-making tools

Processing

- Live 3D reconstruction
- Live drift correction
- Robust, fast, efficient postreconstruction
- State-of-the art toolbox for localization microscopy

Analysis

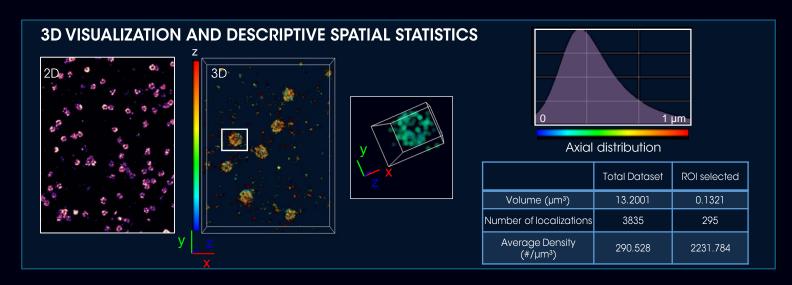
- Complete 3D multi-color viewer solution
- Spatial descriptive statistics
- Density-based spatial clustering
- Resolution analysis
- Single-particle tracking analysis
- Spectral demixing for multicolor imaging





Nanoscopy data, unlike standard microscopy images, are coordinate-based rather than pixel-based, opening up new avenues for in-depth data analysis.

NEO SAFe software offers a variety of tools for nanoscopy data visualization and analysis.



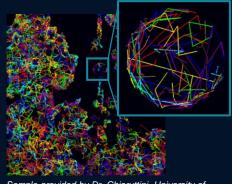
CLUSTERING ANALYSIS Isolation of clusters DBSCAN (Ester et al. 1996) or Voronoi sègmentation (SR-Tesseller) (Levet et al. 2015) Quantification Clusters Centroid 669 678 654 514 525 526 0.313 0.302 0.308 Cluster density (N/µm³) (µm) Number of 277 249 256 ocalizations Volume 0.015 0.019 0.017 (µm³) Cluster volume (um3)

SINGLE PARTICLE TRACKING

Detection and Localization

SPT raw data are similar as standard single-molecule acquisitions, but the movement of each molecule is tracked

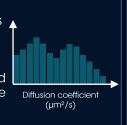
Track reconstruction



Sample provided by Dr. Chiaruttini, University of Geneva, Switzerland

Quantification

- Number of tracks
- Track duration
- Track intensity
- Diffusion
 coefficient based
 on Mean Square
 Displacement
 (MSD)

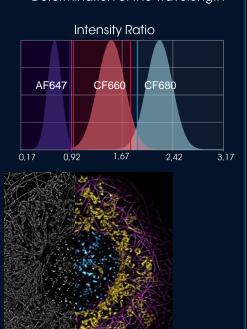


SPECTRAL DEMIXING

Multicolor with far red dyes

For each localization:

- Measurement of the intensity ratio between the 2 cameras
- Determination of the wavelength

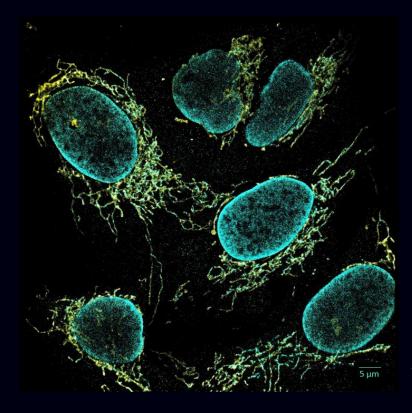




Tubulin – CF 660 Mitochondria – CF 680

EdU - AF 647





DNA in mitochondria



Fixed cells



TOM20-AF 647 dsDNA-CF 568



Colocalization

Prof. Yoerg Bewersdorf Yale University School of Medicine

Axon nanoscale architecture



Rat hippocampal neurons, fixed

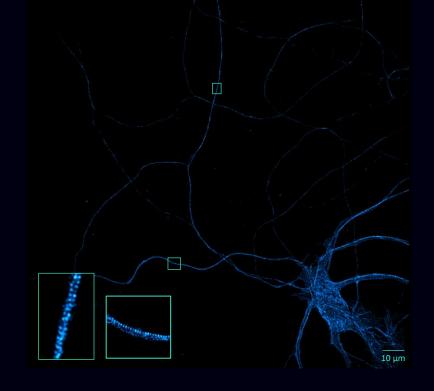


3D-dSTORM β2-spectrin - AF 647

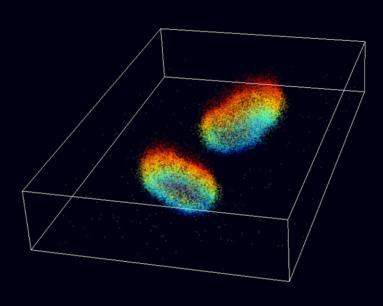


Nano-organisation segmentation

Dr. Christophe Leterrier Neurocytolab, INP Aix-Marseille University







Living bacteria



Escherichia coli, live



dual-color 3D-dSTORM bacterial membrane and wall - AF 647



quantitative 3D nanometric measures

Dr. Clément Cabriel

Sandrine Lévêque-Fort team Institut des Sciences Moléculaires d'Orsay (ISMO)

Emerging model organisms



Trypanosoma brucei, fixed

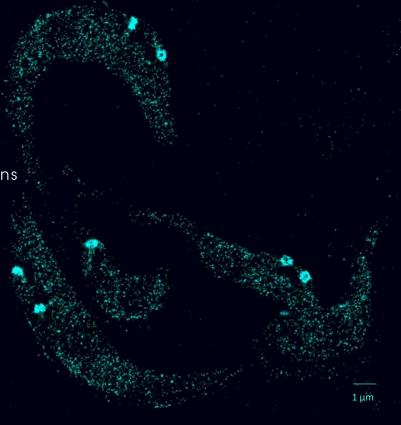


dual-color 3D-dSTORM intra-flagellar transport proteins AF 647



DBSCAN, Voronoi clustering

Dr. Jamin Jung
Trypanosome Cell Biology
Pasteur Institute



abbelight research SAFe hands-on nanoscopy workshop

A 3-full day course on single-molecule localization microscopy combining theory, round tables, practice on SAFe instrument and image analysis on NEO software.

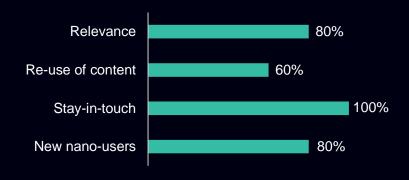
Organized on-site, this workshop is tailored to your needs and adapted to both experts and neophytes to launch as many new projects as possible.

TEST your own samples with abbelight technology

LEARN how to optimize sample preparation, image acquisition

and analysis

LAUNCH your new nanoscopy projects

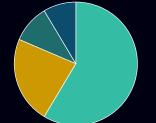


"Very fast and user friendly;
Very nice software, easy to use and easy to manage;
It is a good instrument to see fine details and the nano-organization of the cytoskeleton, of neurons and bacteria, at least this is what we saw live during the workshop. "

Italian Institute of Technology, Genoa 40 participants

"Before the demo, the data analysis and the reconstruction was quite of a "black box" to me (at least the way the software reconstructs the images after the acquisition). Now everything is much clearer. Thanks so much!

Rennes, France 10 participants Will you use 3D nanoscopy in your future research?

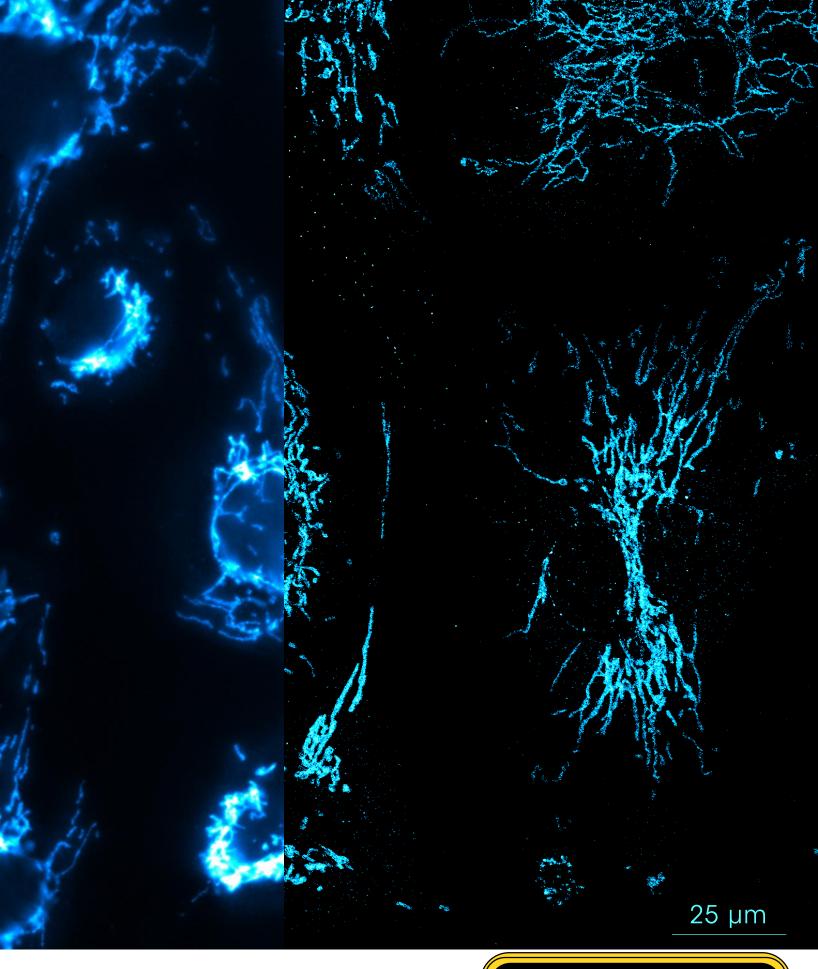


- Yes, in the upcoming months
- Yes, as soon as possible
- I do not know
- No



	and nanoscopy			
	Microscopy market Compilation of the best features of the best optical microscopes	Nanoscopy market Compilation of the best features of the best single-molecule instruments	abbelight SAFe 180	abbelight SAFe 360
Localization precision xyz (nm³)	N.A. (Lateral resolution : 120-150 nm)	10*10*25	10*10*25	10*10*15
Method for 3D	Z- stack	Astigmatism	Astigmatism	Astigmatism & super- critical angle fluorescence
Parameters affecting lateral resolution (other than sample prep)		Illumination heterogeneity, lateral drift	None (homogeneous illumination) & Cross-correlation	None (homogeneous illumination) & Cross-correlation
Field of view (µm²)	150*150	50*50	150*150	150*150
Multicolor imaging	Simultaneous multicolor imaging	Simultaneous multicolor imaging	4 colors sequentially	Simultaneous multicolor imaging (spectral demixing)
Excitation modality	Scanning	Epi, TIRF, Hilo	Epi, TIRF, Hilo	Epi, TIRF, Hilo
Standard imaging modalities	FRAP, FRET	PALM, STORM, SPT, TIRF	PALM, STORM, SPT, TIRF	PALM, STORM, SPT, TIRF
Capture range (µm)	N.A	0,6	0,6	1
Depth of imaging (µm)	> 25	5-10	5-10	5-10
Laser power needed for the largest FOV	< 10 mW	> 1000 mW	< 300 mW	< 300 mW
Acquisition duration	10 ms	Seconds to minutes	Seconds to minutes	Seconds to minutes
Compatibility with existing systems			Confocal, Spinning Disk, Widefield, SIM, STED	Confocal, Spinning Disk, Widefield, SIM, STED
Time to 2D & 3D reconstructed image Drift corrected –largest FOV	Real-time	Real-time	Real-time	Realtime
Live cell imaging	Video Microscopy LSM Spinning disk	Live SIM Live SPT	Video Microscopy (TIRF, EPI, Hilo) Live SPT	Video Microscopy (TIRF, EPI, Hilo) Live SPT
Nanoscale Data analysis		3D visualization SPT analysis Clustering	3D viewer Clustering SPT analysis	3D viewer Clustering SPT analysis Spectral Demixing
Scientific support			SAFe accelerator Abbelight academy - Online tutorials	SAFe accelerator Abbelight academy - Online tutorials
Super-resolution buffer & samples			Smart KIT Blinking Pad Calibration samples	Smart KIT Blinking Pad Calibration samples







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abbelight, 191 avenue Aristide Briand 94230 Cachan, France DANGER-VISIBLE AND INVISIBLE LASER RADIATION AVOID EYE OR SKIN EXPOSURE TO DIRECT OR SCATTERED RADIATION CLASS 4

LASER PRODUCT
Total Power 1500 mW MAX
CW 400-700 nm
IEC/EN60825-1: 2014

Complies with FDA performance standards for laser products except for deviations pursuant to Laser Notice n°50, dated June 24, 2007