

SUPER-RESOLUTION BY STRUCTURED-ILLUMINATION-AXIAL-TOMOGRAPHY (SIAT)

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Key Words: Axial Tomography, SIM, SIAT

Since the first developments of super-resolution-microscopy (SRM), methods allowing to circumvent (“break“) the Abbe limit, a variety of different complementary SRM techniques were implemented. Due to differences in the mode of operation, a further optimization of super-resolution imaging can be achieved by appropriate combinatorial strategies.

One of these is the combination of Structured-Illumination-Microscopy (SIM) and Micro-Axial-Tomography (AT) [1,2]. Both of these microscopy methods operate in the visible range spectrum and have been demonstrated to be excellently suitable for live-cell imaging, due to very low illumination intensities.

While SIM allows for a resolution-improvement of a factor of two along all spatial directions (3D), together with appropriate image processing, SIM-AT is expected to achieve an isotropic effective resolution down to the 100nm region.

We present an experimental realization of the combination of the two techniques together with first results. A major benefit of the SIAT microscope is that it provides super-resolution imaging without challenging conditions on the fluorophores, and that works well also on label-free (autofluorescent) samples.

[1] T. Bruns, S. Schickinger, H. Schneckenburger: „Mikroskopadapter zur Rotation 3-dimensionalen Proben“, *BioPhotonik* (2014) 40-41

[2] R. Heintzmann and C. Cremer, “Axial tomographic confocal fluorescence microscopy”, *J.Microsc.* 206, 7-23 (2002).