Stefan Jakobs

List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/6237364/stefan-jakobs-publications-by-year.pdf

Version: 2024-04-20

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

61 13,039 114 127 h-index g-index citations papers 6.36 138 15,033 10.2 L-index avg, IF ext. papers ext. citations

#	Paper	IF	Citations
127	Optimal precision and accuracy in 4Pi-STORM using dynamic spline PSF models <i>Nature Methods</i> , 2022 , 19, 603-612	21.6	1
126	The TFAM-to-mtDNA ratio defines inner-cellular nucleoid populations with distinct activity levels. <i>Cell Reports</i> , 2021 , 37, 110000	10.6	7
125	MINSTED fluorescence localization and nanoscopy. <i>Nature Photonics</i> , 2021 , 15, 361-366	33.9	18
124	Colocalization for super-resolution microscopy via optimal transport. <i>Nature Computational Science</i> , 2021 , 1, 199-211		2
123	The Positive Switching Fluorescent Protein Padron2 Enables Live-Cell Reversible Saturable Optical Linear Fluorescence Transitions (RESOLFT) Nanoscopy without Sequential Illumination Steps. <i>ACS Nano</i> , 2021 , 15, 9509-9521	16.7	2
122	Correlative fluorescence microscopy, transmission electron microscopy and secondary ion mass spectrometry (CLEM-SIMS) for cellular imaging. <i>PLoS ONE</i> , 2021 , 16, e0240768	3.7	2
121	Monitoring mitochondrial translation in living cells. <i>EMBO Reports</i> , 2021 , 22, e51635	6.5	6
120	Innenarchitektur der Zellkraftwerke [Membranfalten in Hochaufl\(\bar{B}\) ung. BioSpektrum, 2021 , 27, 161-164	0.1	
119	isoSTED microscopy with water-immersion lenses and background reduction. <i>Biophysical Journal</i> , 2021 , 120, 3303-3314	2.9	2
118	Mapping protein interactions in the active TOM-TIM23 supercomplex. <i>Nature Communications</i> , 2021 , 12, 5715	17.4	5
117	Reversibly Switchable Fluorescent Proteins for RESOLFT Nanoscopy. <i>Topics in Applied Physics</i> , 2020 , 247	1-2 6 1	5
116	MICOS assembly controls mitochondrial inner membrane remodeling and crista junction redistribution to mediate cristae formation. <i>EMBO Journal</i> , 2020 , 39, e104105	13	43
115	Light Microscopy of Mitochondria at the Nanoscale. <i>Annual Review of Biophysics</i> , 2020 , 49, 289-308	21.1	21
114	Photoswitching mechanism of a fluorescent protein revealed by time-resolved crystallography and transient absorption spectroscopy. <i>Nature Communications</i> , 2020 , 11, 741	17.4	23
113	Kinetic coupling of the respiratory chain with ATP synthase, but not proton gradients, drives ATP production in cristae membranes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020 , 117, 2412-2421	11.5	25
112	Absolute quantum yield measurements of fluorescent proteins using a plasmonic nanocavity. <i>Communications Biology</i> , 2020 , 3, 627	6.7	4
111	Multicolor 3D MINFLUX nanoscopy of mitochondrial MICOS proteins. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020 , 117, 20607-20614	11.5	25

110	Live-cell RESOLFT nanoscopy of transgenic. <i>Plant Direct</i> , 2020 , 4, e00261	3.3	2
109	Live-cell STED nanoscopy of mitochondrial cristae. <i>Scientific Reports</i> , 2019 , 9, 12419	4.9	75
108	Correlative cryo super-resolution light and electron microscopy on mammalian cells using fluorescent proteins. <i>Scientific Reports</i> , 2019 , 9, 1369	4.9	55
107	Correlative STED super-resolution light and electron microscopy on resin sections. <i>Journal Physics D: Applied Physics</i> , 2019 , 52, 374003	3	4
106	Mitochondrial fusion is required for regulation of mitochondrial DNA replication. <i>PLoS Genetics</i> , 2019 , 15, e1008085	6	62
105	A MICOS-TIM22 Association Promotes Carrier Import into Human Mitochondria. <i>Journal of Molecular Biology</i> , 2019 , 431, 2835-2851	6.5	30
104	Mic60 exhibits a coordinated clustered distribution along and across yeast and mammalian mitochondria. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 9853-9858	11.5	28
103	Molecular contribution function in RESOLFT nanoscopy. <i>Optics Express</i> , 2019 , 27, 21956-21987	3.3	4
102	ROMO1 is a constituent of the human presequence translocase required for YME1L protease import. <i>Journal of Cell Biology</i> , 2019 , 218, 598-614	7.3	24
101	Spatial orchestration of mitochondrial translation and OXPHOS complex assembly. <i>Nature Cell Biology</i> , 2018 , 20, 528-534	23.4	57
100	Aberration-corrected cryoimmersion light microscopy. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018 , 115, 1204-1209	11.5	20
99	Two-Color 810 nm STED Nanoscopy of Living Cells with Endogenous SNAP-Tagged Fusion Proteins. <i>ACS Chemical Biology,</i> 2018 , 13, 475-480	4.9	42
98	Chromophore twisting in the excited state of a photoswitchable fluorescent protein captured by time-resolved serial femtosecond crystallography. <i>Nature Chemistry</i> , 2018 , 10, 31-37	17.6	99
97	Near-infrared STED nanoscopy with an engineered bacterial phytochrome. <i>Nature Communications</i> , 2018 , 9, 4762	17.4	22
96	The MICOS component Mic60 displays a conserved membrane-bending activity that is necessary for normal cristae morphology. <i>Journal of Cell Biology</i> , 2017 , 216, 889-899	7.3	55
95	Achromatic light patterning and improved image reconstruction for parallelized RESOLFT nanoscopy. <i>Scientific Reports</i> , 2017 , 7, 44619	4.9	20
94	Fluorescence nanoscopy in cell biology. <i>Nature Reviews Molecular Cell Biology</i> , 2017 , 18, 685-701	48.7	520
93	Bax assembles into large ring-like structures remodeling the mitochondrial outer membrane in apoptosis. <i>EMBO Journal</i> , 2016 , 35, 402-13	13	195

Coordinate-targeted fluorescence nanoscopy with multiple off states. Nature Photonics, 2016, 10, 122-138.9 92 In vivo super-resolution RESOLFT microscopy of Drosophila melanogaster. ELife, 2016, 5, 8.9 91 Comment on "Extended-resolution structured illumination imaging of endocytic and cytoskeletal 90 33.3 31 dynamics". Science, 2016, 352, 527 Reversibel schaltbare fluoreszierende Proteine fildie Superauflung. BioSpektrum, 2016, 22, 365-367 89 0.1 TIM29 is a subunit of the human carrier translocase required for protein transport. FEBS Letters, 88 3.8 42 2016. 590. 4147-4158 CRISPR/Cas9-mediated endogenous protein tagging for RESOLFT super-resolution microscopy of 87 108 4.9 living human cells. Scientific Reports, 2015, 5, 9592 Primary light-induced reaction steps of reversibly photoswitchable fluorescent protein Padron0.9 86 9 3.4 investigated by femtosecond spectroscopy. Journal of Physical Chemistry B, 2015, 119, 5136-44 Mic10 oligomerizes to bend mitochondrial inner membranes at cristae junctions. Cell Metabolism, 85 24.6 93 **2015**, 21, 756-63 Cross-strand binding of TFAM to a single mtDNA molecule forms the mitochondrial nucleoid. 84 11.5 193 Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 11288-93 Expression-Enhanced Fluorescent Proteins Based on Enhanced Green Fluorescent Protein for 83 16.7 64 Super-resolution Microscopy. ACS Nano, 2015, 9, 9528-41 The Oxidation Status of Mic19 Regulates MICOS Assembly. Molecular and Cellular Biology, 2015, 35, 4222£87 82 MITRAC7 Acts as a COX1-Specific Chaperone and Reveals a Checkpoint during Cytochrome c 81 10.6 38 Oxidase Assembly. *Cell Reports*, **2015**, 12, 1644-55 80 The 2015 super-resolution microscopy roadmap. Journal Physics D: Applied Physics, 2015, 48, 443001 3 211 Organization of Mitochondrial Gene Expression in Two Distinct Ribosome-Containing Assemblies. 10.6 64 79 Cell Reports, 2015, 10, 843-853 RESOLFT Nanoscopy of Fixed Cells Using a Z-Domain Based Fusion Protein for Labelling. PLoS ONE, 78 3.7 4 **2015**, 10, e0136233 Two-color RESOLFT nanoscopy with green and red fluorescent photochromic proteins. 39 77 3.2 ChemPhysChem, 2014, 15, 655-63 Super-resolution microscopy of mitochondria. Current Opinion in Chemical Biology, 2014, 20, 9-15 76 81 9.7 Uniform nomenclature for the mitochondrial contact site and cristae organizing system. Journal of 7.3 177 Cell Biology, 2014, 204, 1083-6

(2011-2014)

74	Live-cell multiplane three-dimensional super-resolution optical fluctuation imaging. <i>Nature Communications</i> , 2014 , 5, 5830	17.4	105
73	Coordinate-targeted and coordinate-stochastic super-resolution microscopy with the reversibly switchable fluorescent protein Dreiklang. <i>ChemPhysChem</i> , 2014 , 15, 756-62	3.2	17
72	STED super-resolution microscopy of clinical paraffin-embedded human rectal cancer tissue. <i>PLoS ONE</i> , 2014 , 9, e101563	3.7	28
71	Nanoscopy with more than 100,000 'doughnuts'. <i>Nature Methods</i> , 2013 , 10, 737-40	21.6	190
70	SNAP-, CLIP- and Halo-tag labelling of budding yeast cells. <i>PLoS ONE</i> , 2013 , 8, e78745	3.7	72
69	STED super-resolution microscopy reveals an array of MINOS clusters along human mitochondria. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013 , 110, 8936-41	11.5	132
68	Nanoscopy of living brain slices with low light levels. <i>Neuron</i> , 2012 , 75, 992-1000	13.9	106
67	Rcf1 mediates cytochrome oxidase assembly and respirasome formation, revealing heterogeneity of the enzyme complex. <i>Cell Metabolism</i> , 2012 , 15, 336-47	24.6	165
66	Red-emitting rhodamines with hydroxylated, sulfonated, and phosphorylated dye residues and their use in fluorescence nanoscopy. <i>Chemistry - A European Journal</i> , 2012 , 18, 12986-98	4.8	36
65	Novel red fluorophores with superior performance in STED microscopy. <i>Optical Nanoscopy</i> , 2012 , 1, 7		68
65 64	Novel red fluorophores with superior performance in STED microscopy. <i>Optical Nanoscopy</i> , 2012 , 1, 7 The inner-mitochondrial distribution of Oxa1 depends on the growth conditions and on the availability of substrates. <i>Molecular Biology of the Cell</i> , 2012 , 23, 2292-301	3.5	68
	The inner-mitochondrial distribution of Oxa1 depends on the growth conditions and on the	3.5	
64	The inner-mitochondrial distribution of Oxa1 depends on the growth conditions and on the availability of substrates. <i>Molecular Biology of the Cell</i> , 2012 , 23, 2292-301 MINOS1 is a conserved component of mitofilin complexes and required for mitochondrial function		26
64	The inner-mitochondrial distribution of Oxa1 depends on the growth conditions and on the availability of substrates. <i>Molecular Biology of the Cell</i> , 2012 , 23, 2292-301 MINOS1 is a conserved component of mitofilin complexes and required for mitochondrial function and cristae organization. <i>Molecular Biology of the Cell</i> , 2012 , 23, 247-57	3.5	26 141
64 63 62	The inner-mitochondrial distribution of Oxa1 depends on the growth conditions and on the availability of substrates. <i>Molecular Biology of the Cell</i> , 2012 , 23, 2292-301 MINOS1 is a conserved component of mitofilin complexes and required for mitochondrial function and cristae organization. <i>Molecular Biology of the Cell</i> , 2012 , 23, 247-57 rsEGFP2 enables fast RESOLFT nanoscopy of living cells. <i>ELife</i> , 2012 , 1, e00248 Nanoscale distribution of mitochondrial import receptor Tom20 is adjusted to cellular conditions and exhibits an inner-cellular gradient. <i>Proceedings of the National Academy of Sciences of the</i>	3.5 8.9	26 141 155
64 63 62 61	The inner-mitochondrial distribution of Oxa1 depends on the growth conditions and on the availability of substrates. <i>Molecular Biology of the Cell</i> , 2012 , 23, 2292-301 MINOS1 is a conserved component of mitofilin complexes and required for mitochondrial function and cristae organization. <i>Molecular Biology of the Cell</i> , 2012 , 23, 247-57 rsEGFP2 enables fast RESOLFT nanoscopy of living cells. <i>ELife</i> , 2012 , 1, e00248 Nanoscale distribution of mitochondrial import receptor Tom20 is adjusted to cellular conditions and exhibits an inner-cellular gradient. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 13546-51	3.5 8.9	26 141 155 100
64 63 62 61 60	The inner-mitochondrial distribution of Oxa1 depends on the growth conditions and on the availability of substrates. <i>Molecular Biology of the Cell</i> , 2012 , 23, 2292-301 MINOS1 is a conserved component of mitofilin complexes and required for mitochondrial function and cristae organization. <i>Molecular Biology of the Cell</i> , 2012 , 23, 247-57 rsEGFP2 enables fast RESOLFT nanoscopy of living cells. <i>ELife</i> , 2012 , 1, e00248 Nanoscale distribution of mitochondrial import receptor Tom20 is adjusted to cellular conditions and exhibits an inner-cellular gradient. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 13546-51 Dual-label STED nanoscopy of living cells using photochromism. <i>Nano Letters</i> , 2011 , 11, 3970-3 A reversibly photoswitchable GFP-like protein with fluorescence excitation decoupled from	3.5 8.9 11.5 11.5	26 141 155 100

56	Super-resolution microscopy reveals that mammalian mitochondrial nucleoids have a uniform size and frequently contain a single copy of mtDNA. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 13534-9	11.5	360
55	Recycling, clustering, and endocytosis jointly maintain PIN auxin carrier polarity at the plasma membrane. <i>Molecular Systems Biology</i> , 2011 , 7, 540	12.2	188
54	Rapid FlAsH labelling in the budding yeast Saccharomyces cerevisiae. <i>Journal of Microscopy</i> , 2010 , 240, 6-13	1.9	8
53	Molecular basis of the light-driven switching of the photochromic fluorescent protein Padron. Journal of Biological Chemistry, 2010 , 285, 14603-9	5.4	58
52	Stimulated emission depletion nanoscopy of living cells using SNAP-tag fusion proteins. <i>Biophysical Journal</i> , 2010 , 98, 158-63	2.9	113
51	Multicolor fluorescence nanoscopy in fixed and living cells by exciting conventional fluorophores with a single wavelength. <i>Biophysical Journal</i> , 2010 , 99, 2686-94	2.9	149
50	Far-field autofluorescence nanoscopy. <i>Nano Letters</i> , 2010 , 10, 4249-52	11.5	21
49	Sample preparation for STED microscopy. <i>Methods in Molecular Biology</i> , 2010 , 591, 185-99	1.4	43
48	New fluorinated rhodamines for optical microscopy and nanoscopy. <i>Chemistry - A European Journal</i> , 2010 , 16, 4477-88	4.8	77
47	Rhodamine NN: eine neue Klasse maskierter Fluoreszenzfarbstoffe. <i>Angewandte Chemie</i> , 2010 , 122, 3598-3602	3.6	31
46	Rhodamines NN: a novel class of caged fluorescent dyes. <i>Angewandte Chemie - International Edition</i> , 2010 , 49, 3520-3	16.4	132
45	Cover Picture: Rhodamines NN: A Novel Class of Caged Fluorescent Dyes (Angew. Chem. Int. Ed. 20/2010). <i>Angewandte Chemie - International Edition</i> , 2010 , 49, 3391-3391	16.4	
44	Two-color STED microscopy reveals different degrees of colocalization between hexokinase-I and the three human VDAC isoforms. <i>PMC Biophysics</i> , 2010 , 3, 4		90
43	The m-AAA protease processes cytochrome c peroxidase preferentially at the inner boundary membrane of mitochondria. <i>Molecular Biology of the Cell</i> , 2009 , 20, 572-80	3.5	30
42	Mitochondrial cristae revealed with focused light. <i>Nano Letters</i> , 2009 , 9, 2508-10	11.5	119
41	Photoswitchable fluorescent proteins enable monochromatic multilabel imaging and dual color fluorescence nanoscopy. <i>Nature Biotechnology</i> , 2008 , 26, 1035-40	44.5	251
40	Spherical nanosized focal spot unravels the interior of cells. <i>Nature Methods</i> , 2008 , 5, 539-44	21.6	323
39	Fluorescence nanoscopy by ground-state depletion and single-molecule return. <i>Nature Methods</i> , 2008 , 5, 943-5	21.6	628

(2005-2008)

38	Nanoscale separation of molecular species based on their rotational mobility. <i>Optics Express</i> , 2008 , 16, 21093-104	3.3	32
37	The class V myosin motor protein, Myo2, plays a major role in mitochondrial motility in Saccharomyces cerevisiae. <i>Journal of Cell Biology</i> , 2008 , 181, 119-30	7.3	90
36	Generation of monomeric reversibly switchable red fluorescent proteins for far-field fluorescence nanoscopy. <i>Biophysical Journal</i> , 2008 , 95, 2989-97	2.9	126
35	Two-color far-field fluorescence nanoscopy. <i>Biophysical Journal</i> , 2007 , 92, L67-9	2.9	205
34	Fluorescence nanoscopy in whole cells by asynchronous localization of photoswitching emitters. <i>Biophysical Journal</i> , 2007 , 93, 3285-90	2.9	227
33	Wide-field subdiffraction RESOLFT microscopy using fluorescent protein photoswitching. <i>Microscopy Research and Technique</i> , 2007 , 70, 269-80	2.8	95
32	Reversible photoswitching enables single-molecule fluorescence fluctuation spectroscopy at high molecular concentration. <i>Microscopy Research and Technique</i> , 2007 , 70, 1003-9	2.8	20
31	Cyclin-dependent kinase 5 is an upstream regulator of mitochondrial fission during neuronal apoptosis. <i>Cell Death and Differentiation</i> , 2007 , 14, 651-61	12.7	91
30	Two-color far-field fluorescence nanoscopy based on photoswitchable emitters. <i>Applied Physics B: Lasers and Optics</i> , 2007 , 88, 161-165	1.9	133
29	1.8 A bright-state structure of the reversibly switchable fluorescent protein Dronpa guides the generation of fast switching variants. <i>Biochemical Journal</i> , 2007 , 402, 35-42	3.8	203
28	Structural basis for reversible photoswitching in Dronpa. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007 , 104, 13005-9	11.5	223
27	High resolution imaging of live mitochondria. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2006 , 1763, 561-75	4.9	99
26	Fis1p and Caf4p, but not Mdv1p, determine the polar localization of Dnm1p clusters on the mitochondrial surface. <i>Journal of Cell Science</i> , 2006 , 119, 3098-106	5.3	60
25	Nanoscale Resolution with Focused Light: Stimulated Emission Depletion and Other Reversible Saturable Optical Fluorescence Transitions Microscopy Concepts 2006 , 571-579		18
24	Differential protein distributions define two sub-compartments of the mitochondrial inner membrane in yeast. <i>FEBS Letters</i> , 2006 , 580, 5628-34	3.8	92
23	4Pi microscopy of quantum dot-labeled cellular structures. <i>Journal of Structural Biology</i> , 2006 , 156, 517	'- <u>3</u> 34	25
22	Nanoscale resolution in GFP-based microscopy. <i>Nature Methods</i> , 2006 , 3, 721-3	21.6	283
21	Breaking the diffraction barrier in fluorescence microscopy at low light intensities by using reversibly photoswitchable proteins. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005 , 102, 17565-9	11.5	632

20	Structure and mechanism of the reversible photoswitch of a fluorescent protein. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005 , 102, 13070-4	11.5	222
19	Mdm31 and Mdm32 are inner membrane proteins required for maintenance of mitochondrial shape and stability of mitochondrial DNA nucleoids in yeast. <i>Journal of Cell Biology</i> , 2005 , 168, 103-15	7.3	88
18	Short tetracysteine tags to beta-tubulin demonstrate the significance of small labels for live cell imaging. <i>Molecular Biology of the Cell</i> , 2004 , 15, 5616-22	3.5	115
17	Concepts for nanoscale resolution in fluorescence microscopy. <i>Current Opinion in Neurobiology</i> , 2004 , 14, 599-609	7.6	226
16	Cooperative 4Pi excitation and detection yields sevenfold sharper optical sections in live-cell microscopy. <i>Biophysical Journal</i> , 2004 , 87, 4146-52	2.9	94
15	Imaging and writing at the nanoscale with focused visible light through saturable optical transitions. <i>Applied Physics A: Materials Science and Processing</i> , 2003 , 77, 859-860	2.6	145
14	Immunofluorescence stimulated emission depletion microscopy. <i>Nature Biotechnology</i> , 2003 , 21, 1303-	4 44.5	126
13	Photoconversion of matrix targeted GFP enables analysis of continuity and intermixing of the mitochondrial lumen. <i>FEBS Letters</i> , 2003 , 554, 194-200	3.8	34
12	Spatial and temporal dynamics of budding yeast mitochondria lacking the division component Fis1p. <i>Journal of Cell Science</i> , 2003 , 116, 2005-14	5.3	82
11	The inner membrane protein Mdm33 controls mitochondrial morphology in yeast. <i>Journal of Cell Biology</i> , 2003 , 160, 553-64	7-3	101
10	Fast 100-nm resolution three-dimensional microscope reveals structural plasticity of mitochondria in live yeast. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2002 , 99, 3370-5	11.5	241
9	Dual-color 4Pi-confocal microscopy with 3D-resolution in the 100 nm range. <i>Ultramicroscopy</i> , 2001 , 90, 207-13	3.1	26
8	4Pi-confocal microscopy of live cells. <i>Ultramicroscopy</i> , 2001 , 87, 155-64	3.1	50
7	Fluorescence microscopy with diffraction resolution barrier broken by stimulated emission. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 8206-10	11.5	1286
6	EFGP and DsRed expressing cultures of Escherichia coli imaged by confocal, two-photon and fluorescence lifetime microscopy. <i>FEBS Letters</i> , 2000 , 479, 131-5	3.8	136
5	Ultrafast dynamics microscopy. <i>Applied Physics Letters</i> , 2000 , 77, 597-599	3.4	31
4	ElAsHIProtein Labeling73-88		1
3	Live-cell STED microscopy of mitochondrial cristae		1

The Positive Switching RSFP Padron2 Enables Live-Cell RESOLFT Nanoscopy Without Sequential Irradiation Steps

1

MINSTED fluorescence localization and nanoscopy

3