

Martin Kampmann

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4224719/publications.pdf>

Version: 2024-02-01

72
papers

9,245
citations

76196

40
h-index

95083

68
g-index

109
all docs

109
docs citations

109
times ranked

14074
citing authors

#	ARTICLE	IF	CITATIONS
1	Genome-Scale CRISPR-Mediated Control of Gene Repression and Activation. <i>Cell</i> , 2014, 159, 647-661.	13.5	2,176
2	Compact and highly active next-generation libraries for CRISPR-mediated gene repression and activation. <i>ELife</i> , 2016, 5, .	2.8	609
3	Mitochondrial stress is relayed to the cytosol by an OMA1-DELE1-HRI pathway. <i>Nature</i> , 2020, 579, 427-432.	13.7	343
4	A Systematic Mammalian Genetic Interaction Map Reveals Pathways Underlying Ricin Susceptibility. <i>Cell</i> , 2013, 152, 909-922.	13.5	332
5	LRP1 is a master regulator of tau uptake and spread. <i>Nature</i> , 2020, 580, 381-385.	13.7	326
6	Targeting the AAA ATPase p97 as an Approach to Treat Cancer through Disruption of Protein Homeostasis. <i>Cancer Cell</i> , 2015, 28, 653-665.	7.7	319
7	Genome-wide programmable transcriptional memory by CRISPR-based epigenome editing. <i>Cell</i> , 2021, 184, 2503-2519.e17.	13.5	312
8	CRISPR Interference-Based Platform for Multimodal Genetic Screens in Human iPSC-Derived Neurons. <i>Neuron</i> , 2019, 104, 239-255.e12.	3.8	288
9	CRISPRi and CRISPRa Screens in Mammalian Cells for Precision Biology and Medicine. <i>ACS Chemical Biology</i> , 2018, 13, 406-416.	1.6	248
10	Molecular characterization of selectively vulnerable neurons in Alzheimer's disease. <i>Nature Neuroscience</i> , 2021, 24, 276-287.	7.1	238
11	Mapping the Genetic Landscape of Human Cells. <i>Cell</i> , 2018, 174, 953-967.e22.	13.5	226
12	Pharmacological dimerization and activation of the exchange factor eIF2B antagonizes the integrated stress response. <i>ELife</i> , 2015, 4, e07314.	2.8	212
13	Combined CRISPRi/a-Based Chemical Genetic Screens Reveal that Rigosertib Is a Microtubule-Destabilizing Agent. <i>Molecular Cell</i> , 2017, 68, 210-223.e6.	4.5	197
14	Heterochromatin anomalies and double-stranded RNA accumulation underlie <i>C9orf72</i> poly(PR) toxicity. <i>Science</i> , 2019, 363, .	6.0	181
15	The mTOR Complex Controls HIV Latency. <i>Cell Host and Microbe</i> , 2016, 20, 785-797.	5.1	179
16	Probing the Global Cellular Responses to Lipotoxicity Caused by Saturated Fatty Acids. <i>Molecular Cell</i> , 2019, 74, 32-44.e8.	4.5	170
17	Genome-wide CRISPRi/a screens in human neurons link lysosomal failure to ferroptosis. <i>Nature Neuroscience</i> , 2021, 24, 1020-1034.	7.1	170
18	Tau Internalization is Regulated by 6-O Sulfation on Heparan Sulfate Proteoglycans (HSPGs). <i>Scientific Reports</i> , 2018, 8, 6382.	1.6	162

#	ARTICLE	IF	CITATIONS
19	Parallel shRNA and CRISPR-Cas9 screens enable antiviral drug target identification. <i>Nature Chemical Biology</i> , 2016, 12, 361-366.	3.9	157
20	Integrated platform for genome-wide screening and construction of high-density genetic interaction maps in mammalian cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, E2317-26.	3.3	121
21	Dual gene activation and knockout screen reveals directional dependencies in genetic networks. <i>Nature Biotechnology</i> , 2018, 36, 170-178.	9.4	120
22	Three-dimensional structure and flexibility of a membrane-coating module of the nuclear pore complex. <i>Nature Structural and Molecular Biology</i> , 2009, 16, 782-788.	3.6	113
23	The Psychiatric Cell Map Initiative: A Convergent Systems Biological Approach to Illuminating Key Molecular Pathways in Neuropsychiatric Disorders. <i>Cell</i> , 2018, 174, 505-520.	13.5	108
24	Validation of the Hsp70- β 3 Protein-Protein Interaction as a Potential Therapeutic Target in Cancer. <i>Molecular Cancer Therapeutics</i> , 2015, 14, 642-648.	1.9	105
25	Compromised function of the ESCRT pathway promotes endolysosomal escape of tau seeds and propagation of tau aggregation. <i>Journal of Biological Chemistry</i> , 2019, 294, 18952-18966.	1.6	103
26	CRISPR-based functional genomics for neurological disease. <i>Nature Reviews Neurology</i> , 2020, 16, 465-480.	4.9	89
27	Paradoxical resistance of multiple myeloma to proteasome inhibitors by decreased levels of 19S proteasomal subunits. <i>ELife</i> , 2015, 4, e08153.	2.8	84
28	Next-generation libraries for robust RNA interference-based genome-wide screens. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, E3384-91.	3.3	83
29	Mapping the orientation of nuclear pore proteins in living cells with polarized fluorescence microscopy. <i>Nature Structural and Molecular Biology</i> , 2011, 18, 643-649.	3.6	81
30	Functional genomics platform for pooled screening and generation of mammalian genetic interaction maps. <i>Nature Protocols</i> , 2014, 9, 1825-1847.	5.5	79
31	Next-Generation NAMPT Inhibitors Identified by Sequential High-Throughput Phenotypic Chemical and Functional Genomic Screens. <i>Chemistry and Biology</i> , 2013, 20, 1352-1363.	6.2	72
32	Reverse gyrase has heat-protective DNA chaperone activity independent of supercoiling. <i>Nucleic Acids Research</i> , 2004, 32, 3537-3545.	6.5	70
33	Gene expression and cell identity controlled by anaphase-promoting complex. <i>Nature</i> , 2020, 579, 136-140.	13.7	69
34	Deep mutational scanning reveals the structural basis for α -synuclein activity. <i>Nature Chemical Biology</i> , 2020, 16, 653-659.	3.9	67
35	A Comprehensive Resource for Induced Pluripotent Stem Cells from Patients with Primary Tauopathies. <i>Stem Cell Reports</i> , 2019, 13, 939-955.	2.3	62
36	Structure of a trimeric nucleoporin complex reveals alternate oligomerization states. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 17693-17698.	3.3	57

#	ARTICLE	IF	CITATIONS
37	Fluorescence Anisotropy Reveals Order and Disorder of Protein Domains in the Nuclear Pore Complex. <i>Biophysical Journal</i> , 2010, 99, 1706-1717.	0.2	54
38	Suppression of B-cell development genes is key to glucocorticoid efficacy in treatment of acute lymphoblastic leukemia. <i>Blood</i> , 2017, 129, 3000-3008.	0.6	48
39	Image-based pooled whole-genome CRISPRi screening for subcellular phenotypes. <i>Journal of Cell Biology</i> , 2021, 220, .	2.3	48
40	A high-throughput screen of real-time ATP levels in individual cells reveals mechanisms of energy failure. <i>PLoS Biology</i> , 2018, 16, e2004624.	2.6	47
41	BRD2 inhibition blocks SARS-CoV-2 infection by reducing transcription of the host cell receptor ACE2. <i>Nature Cell Biology</i> , 2022, 24, 24-34.	4.6	47
42	Ceapins block the unfolded protein response sensor ATF6 β by inducing a neomorphic inter-organelle tether. <i>ELife</i> , 2019, 8, .	2.8	46
43	Unraveling the mechanism of cell death induced by chemical fibrils. <i>Nature Chemical Biology</i> , 2014, 10, 969-976.	3.9	43
44	A CRISPR Approach to Neurodegenerative Diseases. <i>Trends in Molecular Medicine</i> , 2017, 23, 483-485.	3.5	41
45	CRISPR-based screens uncover determinants of immunotherapy response in multiple myeloma. <i>Blood Advances</i> , 2020, 4, 2899-2911.	2.5	36
46	Knocking out the door to tunicamycin entry. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 11731-11732.	3.3	32
47	Relapse-associated AURKB blunts the glucocorticoid sensitivity of B cell acute lymphoblastic leukemia. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 3052-3061.	3.3	32
48	Obstacle Bypass in Protein Motion along DNA by Two-dimensional Rather than One-dimensional Sliding. <i>Journal of Biological Chemistry</i> , 2004, 279, 38715-38720.	1.6	31
49	Conserved Spatial Organization of FG Domains in the Nuclear Pore Complex. <i>Biophysical Journal</i> , 2013, 104, 37-50.	0.2	31
50	Weak base pairing in both seed and 3'UTR regions reduces RNAi off-targets and enhances si/shRNA designs. <i>Nucleic Acids Research</i> , 2014, 42, 12169-12176.	6.5	27
51	Elucidating drug targets and mechanisms of action by genetic screens in mammalian cells. <i>Chemical Communications</i> , 2017, 53, 7162-7167.	2.2	26
52	Pharmaceutical-Grade Rigosertib Is a Microtubule-Destabilizing Agent. <i>Molecular Cell</i> , 2020, 79, 191-198.e3.	4.5	22
53	Facilitated diffusion in chromatin lattices: mechanistic diversity and regulatory potential. <i>Molecular Microbiology</i> , 2005, 57, 889-899.	1.2	20
54	Genome-wide CRISPRi screening identifies OCIAD1 as a prohibitin client and regulatory determinant of mitochondrial Complex III assembly in human cells. <i>ELife</i> , 2021, 10, .	2.8	20

#	ARTICLE	IF	CITATIONS
55	NudC guides client transfer between the Hsp40/70 and Hsp90 chaperone systems. <i>Molecular Cell</i> , 2022, 82, 555-569.e7.	4.5	20
56	CRISPulator: a discrete simulation tool for pooled genetic screens. <i>BMC Bioinformatics</i> , 2017, 18, 347.	1.2	19
57	Extending chemical perturbations of the ubiquitin fitness landscape in a classroom setting reveals new constraints on sequence tolerance. <i>Biology Open</i> , 2018, 7, .	0.6	17
58	Defining the ATPome reveals cross-optimization of metabolic pathways. <i>Nature Communications</i> , 2020, 11, 4319.	5.8	17
59	A high-throughput CRISPR interference screen for dissecting functional regulators of GPCR/cAMP signaling. <i>PLoS Genetics</i> , 2020, 16, e1009103.	1.5	15
60	Use of RNA Tertiary Interaction Modules for the Crystallisation of the Spliceosomal snRNP Core Domain. <i>Journal of Molecular Biology</i> , 2010, 402, 154-164.	2.0	11
61	Robust Sequence Determinants of $\hat{\pm}$ -Synuclein Toxicity in Yeast Implicate Membrane Binding. <i>ACS Chemical Biology</i> , 2020, 15, 2137-2153.	1.6	9
62	CRISPR-based genetic interaction maps inform therapeutic strategies in cancer. <i>Translational Cancer Research</i> , 2018, 7, S61-S67.	0.4	8
63	Patterns of neuronal Rhes as a novel hallmark of tauopathies. <i>Acta Neuropathologica</i> , 2021, 141, 651-666.	3.9	6
64	Selective vulnerabilities in the proteostasis network of castration-resistant prostate cancer. <i>Cell Chemical Biology</i> , 2022, 29, 490-501.e4.	2.5	6
65	Functional genomics screen identifies proteostasis targets that modulate prion protein (PrP) stability. <i>Cell Stress and Chaperones</i> , 2021, 26, 443-452.	1.2	5
66	Nascent Proteins Caught in the Act. <i>Science</i> , 2009, 326, 1352-1353.	6.0	4
67	Phenotypic Screening Using High-Content Imaging to Identify Lysosomal pH Modulators in a Neuronal Cell Model. <i>ACS Chemical Neuroscience</i> , 2022, , .	1.7	3
68	A high-throughput CRISPR interference screen for dissecting functional regulators of GPCR/cAMP signaling. <i>FASEB Journal</i> , 2021, 35, .	0.2	1
69	Conversations with LGBT+ scientists about visibility, leadership and climbing the career ladder. <i>Journal of Cell Science</i> , 2022, 135, .	1.2	1
70	Pre-Clinical Activity of the Novel, First-in-Class p97 Inhibitor, CB-5083, in Multiple Myeloma. <i>Blood</i> , 2014, 124, 4701-4701.	0.6	0
71	Functional Multi-Omics Reveals Genetic and Pharmacologic Regulation of Surface CD38 in Multiple Myeloma. <i>Blood</i> , 2021, 138, 2648-2648.	0.6	0
72	CRISPR-Based Screening for Stress Response Factors in Mammalian Cells. <i>Methods in Molecular Biology</i> , 2022, 2428, 19-40.	0.4	0