

Christoph WÃ¼lfing

List of Publications by Year in descending order

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docs citations

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times ranked

3332
citing authors

#	ARTICLE	IF	CITATIONS
1	Cellular Structures Controlling T Cell Signaling in Time and Space. , 2022, , .		0
2	Adenosine 2A receptor and TIM3 suppress cytolytic killing of tumor cells via cytoskeletal polarization. Communications Biology, 2022, 5, 9.	4.4	4
3	A LAT-Based Signaling Complex in the Immunological Synapse as Determined with Live Cell Imaging Is Less Stable in T Cells with Regulatory Capability. Cells, 2021, 10, 418.	4.1	0
4	PD-1 suppresses the maintenance of cell couples between cytotoxic T cells and target tumor cells within the tumor. Science Signaling, 2020, 13, .	3.6	15
5	Super-resolution Imaging of the T cell Central Supramolecular Signaling Cluster Using Stimulated Emission Depletion Microscopy. Bio-protocol, 2020, 10, e3806.	0.4	1
6	Transient protein accumulation at the center of the T cell antigen-presenting cell interface drives efficient IL-2 secretion. ELife, 2019, 8, .	6.0	7
7	Systems Imaging of the Immune Synapse. Methods in Molecular Biology, 2017, 1584, 409-421.	0.9	9
8	Image-based spatiotemporal causality inference for protein signaling networks. Bioinformatics, 2017, 33, i217-i224.	4.1	1
9	PKC δ links proximal T cell and Notch signaling through localized regulation of the actin cytoskeleton. ELife, 2017, 6, .	6.0	18
10	Computational spatiotemporal analysis identifies WAVE2 and cofilin as joint regulators of costimulation-mediated T cell actin dynamics. Science Signaling, 2016, 9, rs3.	3.6	24
11	Inhibition of diacylglycerol kinase δ restores restimulation-induced cell death and reduces immunopathology in XLP-1. Science Translational Medicine, 2016, 8, 321ra7.	12.4	41
12	Modest Interference with Actin Dynamics in Primary T Cell Activation by Antigen Presenting Cells Preferentially Affects Lamellar Signaling. PLoS ONE, 2015, 10, e0133231.	2.5	8
13	Early Signaling in Primary T Cells Activated by Antigen Presenting Cells Is Associated with a Deep and Transient Lamellar Actin Network. PLoS ONE, 2015, 10, e0133299.	2.5	19
14	T/B δ cell interactions are more transient in response to weak stimuli in SLE δ prone mice. European Journal of Immunology, 2014, 44, 3522-3531.	2.9	18
15	New inhibitory signaling by CTLA-4. Nature Immunology, 2014, 15, 408-409.	14.5	20
16	The lymphoid lineage δ specific actin-uncapping protein Rltpr is essential for costimulation via CD28 and the development of regulatory T cells. Nature Immunology, 2013, 14, 858-866.	14.5	100
17	The actin δ driven spatiotemporal organization of T δ cell signaling at the system scale. Immunological Reviews, 2013, 256, 133-147.	6.0	27
18	New TACTICS for finding Numb. Immunology and Cell Biology, 2013, 91, 1-2.	2.3	2

#	ARTICLE	IF	CITATIONS
19	GRB2-Mediated Recruitment of THEMIS to LAT Is Essential for Thymocyte Development. <i>Journal of Immunology</i> , 2013, 190, 3749-3756.	0.8	71
20	Mechanism and function of Vav1 localization in TCR signaling. <i>Journal of Cell Science</i> , 2012, 125, 5302-14.	2.0	26
21	Phosphatidylinositol (4,5) Bisphosphate Controls T Cell Activation by Regulating T Cell Rigidity and Organization. <i>PLoS ONE</i> , 2011, 6, e27227.	2.5	31
22	Itk Controls the Spatiotemporal Organization of T Cell Activation. <i>Science Signaling</i> , 2011, 4, ra66.	3.6	48
23	The CD3 ζ Subunit Contains a Phosphoinositide-Binding Motif That Is Required for the Stable Accumulation of TCR α CD3 Complex at the Immunological Synapse. <i>Journal of Immunology</i> , 2011, 186, 6839-6847.	0.8	73
24	Inhibiting the Inhibitor of the Inhibitor: Blocking PKC ζ to Enhance Regulatory T Cell Function. <i>Science Signaling</i> , 2010, 3, pe24.	3.6	19
25	Tentative and transient natural killer cell polarization balances the requirements for discriminatory recognition and cytolytic efficacy. <i>Communicative and Integrative Biology</i> , 2010, 3, 545-548.	1.4	3
26	Transience in polarization of cytolytic effectors is required for efficient killing and controlled by Cdc42. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 11912-11917.	7.1	27
27	Spatiotemporal Patterning During T Cell Activation Is Highly Diverse. <i>Science Signaling</i> , 2009, 2, ra15.	3.6	88
28	The Cytoplasmic Tail of the T Cell Receptor CD3 μ Subunit Contains a Phospholipid-Binding Motif that Regulates T Cell Functions. <i>Journal of Immunology</i> , 2009, 183, 1055-1064.	0.8	73
29	Cdc42/Rac function in NK cells and CTLs is variable and governed by spatiotemporal patterning of Cdc42/Rac. <i>FASEB Journal</i> , 2008, 22, 1064.15.	0.5	0
30	Itk regulates T cell signaling through localization of active Cdc42. <i>FASEB Journal</i> , 2008, 22, 1064.18.	0.5	0
31	Protein transduction as a means of effective manipulation of Cdc42 activity in primary T cells. <i>Journal of Immunological Methods</i> , 2007, 319, 64-78.	1.4	9
32	Requirement of homotypic NK-cell interactions through 2B4(CD244)/CD48 in the generation of NK effector functions. <i>Blood</i> , 2006, 107, 3181-3188.	1.4	78
33	A Large T Cell Invagination with CD2 Enrichment Resets Receptor Engagement in the Immunological Synapse. <i>Journal of Immunology</i> , 2006, 177, 4402-4413.	0.8	34
34	Specific Patterns of Cdc42 Activity Are Related to Distinct Elements of T Cell Polarization. <i>Journal of Immunology</i> , 2006, 177, 1708-1720.	0.8	52
35	The CD3 ζ μ / ζ μ signaling module provides normal T α β cell functions in the absence of the TCR α ζ immunoreceptor tyrosine-based activation motifs. <i>European Journal of Immunology</i> , 2005, 35, 3643-3654.	2.9	29
36	T cell receptor (TCR) clustering in the immunological synapse integrates TCR and costimulatory signaling in selected T cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 2904-2909.	7.1	87

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37	Polycomb Group Protein Ezh2 Controls Actin Polymerization and Cell Signaling. <i>Cell</i> , 2005, 121, 425-436.	28.9	345
38	The Murine NK Receptor 2B4 (CD244) Exhibits Inhibitory Function Independent of Signaling Lymphocytic Activation Molecule-Associated Protein Expression. <i>Journal of Immunology</i> , 2004, 173, 3953-3961.	0.8	46
39	Regulation of Sustained Actin Dynamics by the TCR and Costimulation as a Mechanism of Receptor Localization. <i>Journal of Immunology</i> , 2003, 171, 2287-2295.	0.8	91
40	Stepwise cytoskeletal polarization as a series of checkpoints in innate but not adaptive cytolytic killing. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003, 100, 7767-7772.	7.1	104
41	Differential Segregation in a Cell-Cell Contact Interface: The Dynamics of the Immunological Synapse. <i>Biophysical Journal</i> , 2002, 83, 1784-1796.	0.5	101
42	Interface accumulation of receptor/ligand couples in lymphocyte activation: methods, mechanisms, and significance. <i>Immunological Reviews</i> , 2002, 189, 64-83.	6.0	25
43	Neuropilin-1: another neuronal molecule in the "immunological synapse". <i>Nature Immunology</i> , 2002, 3, 418-419.	14.5	9
44	Costimulation and endogenous MHC ligands contribute to T cell recognition. <i>Nature Immunology</i> , 2002, 3, 42-47.	14.5	285
45	Determination of the Relationship Between T Cell Responsiveness and the Number of MHC-Peptide Complexes Using Specific Monoclonal Antibodies. <i>Journal of Immunology</i> , 2000, 164, 5626-5634.	0.8	84
46	Thirty-six views of T cell recognition. <i>Philosophical Transactions of the Royal Society B: Biological Sciences</i> , 2000, 355, 1071-1076.	4.0	25
47	Visualizing lymphocyte recognition. <i>Immunology and Cell Biology</i> , 1999, 77, 186-187.	2.3	6
48	Kinetics and Extent of T Cell Activation as Measured with the Calcium Signal. <i>Journal of Experimental Medicine</i> , 1997, 185, 1815-1825.	8.5	161
49	Altered T Cell Receptor Ligands Trigger a Subset of Early T Cell Signals. <i>Immunity</i> , 1996, 5, 125-135.	14.3	155
50	Protein folding in the periplasm of <i>Escherichia coli</i> . <i>Molecular Microbiology</i> , 1994, 12, 685-692.	2.5	177
51	Correctly Folded T-cell Receptor Fragments in the Periplasm of <i>Escherichia coli</i> . <i>Journal of Molecular Biology</i> , 1994, 242, 655-669.	4.2	83
52	A versatile and highly repressible <i>Escherichia coli</i> expression system based on invertible promoters: expression of a gene encoding a toxic product. <i>Gene</i> , 1993, 136, 199-203.	2.2	18