Satoru Yuzawa

List of Publications by Year in descending order

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623734 752698 23 951 14 20 h-index citations g-index papers 23 23 23 1285 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Intramolecular interaction in LGN, an adaptor protein that regulates mitotic spindle orientation. Journal of Biological Chemistry, 2019, 294, 19655-19666.	3.4	10
2	Ric-8A-mediated stabilization of the trimeric G protein subunit Gαi is inhibited by pertussis toxin-catalyzed ADP-ribosylation. Biochemical and Biophysical Research Communications, 2017, 483, 941-945.	2.1	0
3	Structural basis of cofactor-mediated stabilization and substrate recognition of the $\hat{l}\pm$ -tubulin acetyltransferase $\hat{l}\pm$ TAT1. Biochemical Journal, 2015, 467, 103-113.	3.7	5
4	Structural basis for the recognition of the scaffold protein Frmpd4/Preso1 by the TPR domain of the adaptor protein LGN. Acta Crystallographica Section F, Structural Biology Communications, 2015, 71, 175-183.	0.8	8
5	Ubiquitination of the heterotrimeric G protein $\hat{l}\pm$ subunits $\hat{Gl}\pm\hat{l}$ 2 and $\hat{Gl}\pm\hat{l}$ 4 is prevented by the guanine nucleotide exchange factor Ric-8A. Biochemical and Biophysical Research Communications, 2013, 435, 414-419.	2.1	17
6	Recognition Mechanism of the Cell Polarity Protein Mammalian Inscuteable-LGN Complex. Nihon Kessho Gakkaishi, 2012, 54, 206-212.	0.0	0
7	Structural basis for interaction between the conserved cell polarity proteins Inscuteable and Leu-Gly-Asn repeat-enriched protein (LGN). Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 19210-19215.	7.1	62
8	A Conserved Region between the TPR and Activation Domains of p67 Participates in Activation of the Phagocyte NADPH Oxidase. Journal of Biological Chemistry, 2010, 285, 31435-31445.	3.4	26
9	The Domain Organization of p67 ^{phox} , a Protein Required for Activation of the Superoxide-Producing NADPH Oxidase in Phagocytes. Journal of Innate Immunity, 2009, 1, 543-555.	3.8	23
10	Solution structure of the Grb2 SH2 domain complexed with a high-affinity inhibitor. Journal of Biomolecular NMR, 2008, 42, 197-207.	2.8	20
11	Contacts between membrane proximal regions of the PDGF receptor ectodomain are required for receptor activation but not for receptor dimerization. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 7681-7686.	7.1	71
12	Structural Basis for Activation of the Receptor Tyrosine Kinase KIT by Stem Cell Factor. Cell, 2007, 130, 323-334.	28.9	290
13	Full-length p40phox structure suggests a basis for regulation mechanism of its membrane binding. EMBO Journal, 2007, 26, 1176-1186.	7.8	60
14	Crystallization and preliminary crystallographic analysis of p40phox, a regulatory subunit of NADPH oxidase. Acta Crystallographica Section F: Structural Biology Communications, 2006, 62, 1018-1020.	0.7	3
15	NMR Solution Structure of the Tandem Src Homology 3 Domains of p47 Complexed with a p22 -derived Proline-rich Peptide. Journal of Biological Chemistry, 2006, 281, 3660-3668.	3.4	72
16	Solution Structure of the Tandem Src Homology 3 Domains of p47 in an Autoinhibited Form. Journal of Biological Chemistry, 2004, 279, 29752-29760.	3.4	51
17	Binding of FAD to Cytochrome b558 Is Facilitated during Activation of the Phagocyte NADPH Oxidase, Leading to Superoxide Production. Journal of Biological Chemistry, 2004, 279, 26378-26386.	3.4	33
18	A molecular mechanism for autoinhibition of the tandem SH3 domains of p47phox, the regulatory subunit of the phagocyte NADPH oxidase. Genes To Cells, 2004, 9, 443-456.	1.2	63

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#	Article	IF	CITATIONS
19	Letter to the Editor: Sequence-specific Resonance Assignments of the Tandem SH3 Domains in an Autoinhibitory form of p47phox. Journal of Biomolecular NMR, 2004, 29, 451-452.	2.8	0
20	Backbone assignments of Grb2 complexed with ligand peptides for SH3 and SH2 domains. Journal of Biomolecular NMR, 2003, 27, 185-186.	2.8	3
21	Crystallization and preliminary crystallographic analysis of the autoinhibited form of the tandem SH3 domain of p47phox. Acta Crystallographica Section D: Biological Crystallography, 2003, 59, 1479-1480.	2.5	12
22	Solution structure of Grb2 reveals extensive flexibility necessary for target recognition. Journal of Molecular Biology, 2001, 306, 527-537.	4.2	59
23	Solution structure of the SH2 domain of Grb2 complexed with the Shc-derived phosphotyrosine-containing peptide. Journal of Molecular Biology, 1999, 289, 439-445.	4.2	63