

Kazuki Saito

List of Publications by Year in descending order

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53
papers

2,358
citations

430874

18
h-index

214800

47
g-index

53
all docs

53
docs citations

53
times ranked

2840
citing authors

#	ARTICLE	IF	CITATIONS
1	Solution structure of the zinc finger domain of human RNF144A ubiquitin ligase. <i>Protein Science</i> , 2020, 29, 1836-1842.	7.6	2
2	Zinc finger domain of the human DTX protein adopts a unique RING fold. <i>Protein Science</i> , 2019, 28, 1151-1156.	7.6	6
3	Unique RING finger structure from the human HRD1 protein. <i>Protein Science</i> , 2019, 28, 448-453.	7.6	3
4	Concise machinery for monitoring ubiquitination activities using novel artificial RING fingers. <i>Protein Science</i> , 2018, 27, 1354-1363.	7.6	7
5	Solution structure of the PHD finger from the human KIAA1045 protein. <i>Protein Science</i> , 2018, 27, 987-992.	7.6	1
6	Molecular Design of Artificial Ring Fingers for Detecting Ubiquitination Activities. <i>Biophysical Journal</i> , 2018, 114, 408a.	0.5	0
7	Unique auto-ubiquitination activities of artificial RING fingers in cancer cells. <i>Protein Science</i> , 2018, 27, 1704-1709.	7.6	5
8	Design of a System for Monitoring Ubiquitination Activities of E2 Enzymes Using Engineered RING Finger Proteins. <i>Methods in Molecular Biology</i> , 2018, 1867, 75-87.	0.9	0
9	Highly sensitive detection of E2 activity in ubiquitination using an artificial RING finger. <i>Journal of Peptide Science</i> , 2017, 23, 222-227.	1.4	10
10	The zinc finger domain of RING finger protein 141 reveals a unique RING fold. <i>Protein Science</i> , 2017, 26, 1681-1686.	7.6	5
11	The unique N-terminal zinc finger of synaptotagmin-like protein 4 reveals FYVE structure. <i>Protein Science</i> , 2017, 26, 2451-2457.	7.6	4
12	Development of Molecular Design Strategy of Artificial Ubiquitin-ligases — Toward Cancer Diagnosis Based on Ubiquitination Activities —. <i>Bunseki Kagaku</i> , 2017, 66, 393-402.	0.2	0
13	Prothoracicotropic Hormone. , 2016, , 407-e55-2.		1
14	Ligand-dependent responses of the silkworm prothoracicotropic hormone receptor, Torso, are maintained by unusual intermolecular disulfide bridges in the transmembrane region. <i>Scientific Reports</i> , 2016, 6, 22437.	3.3	5
15	Application of plug-plug technique to <sc>ACE</sc> experiments for discovery of peptides binding to a larger target protein: A model study of calmodulin-binding fragments selected from a digested mixture of reduced <sc>BSA</sc>. <i>Electrophoresis</i> , 2014, 35, 846-854.	2.4	6
16	Application of plug-plug ACE method to drug discovery in the proteomic era. <i>Seibutsu Butsuri Kagaku</i> , 2014, 58, 71-73.	0.1	0
17	Simultaneous quantification of individual intermediate steroids in silkworm ecdysone biosynthesis by liquid chromatography-tandem mass spectrometry with multiple reaction monitoring. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2013, 915-916, 52-56.	2.3	40
18	Molecular basis of wing coloration in a Batesian mimic butterfly, <i>Papilio polytes</i> . <i>Scientific Reports</i> , 2013, 3, 3184.	3.3	44

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19	Evaluation of dimerization-inhibitory activities of cyclic peptides containing a β -hairpin loop sequence of the EGF receptor. <i>Bioorganic and Medicinal Chemistry</i> , 2012, 20, 5730-5737.	3.0	17
20	Intact-cell-based surface plasmon resonance measurements for ligand affinity evaluation of a membrane receptor. <i>Analytical Biochemistry</i> , 2012, 420, 185-187.	2.4	13
21	Quantitative evaluation of refolding conditions for a disulfide-bond-containing protein using a concise ^{18}O -labeling technique. <i>Protein Science</i> , 2011, 20, 1090-1096.	7.6	1
22	Verification of protein disulfide bond arrangement by in-gel tryptic digestion under entirely neutral pH conditions. <i>Proteomics</i> , 2010, 10, 1505-1509.	2.2	7
23	Inhibitory effect of a dimerization-arm-mimetic peptide on EGF receptor activation. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009, 19, 3279-3282.	2.2	17
24	Fmoc-based solid phase chemical synthesis of 71-mer neuregulin β 1, an epidermal growth factor-like domain. <i>Journal of Peptide Science</i> , 2008, 14, 261-266.	1.4	0
25	Novel non-peptidic and small-sized BACE1 inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 1654-1658.	2.2	46
26	BACE1 inhibitors: Optimization by replacing the α -hydroxy acid residue with non-acidic moiety. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 1649-1653.	2.2	40
27	NTA-mediated protein capturing strategy in screening experiments for small organic molecules by surface plasmon resonance. <i>Proteomics</i> , 2007, 7, 494-499.	2.2	9
28	Design and synthesis of potent β -secretase (BACE1) inhibitors with carboxylic acid bioisosteres. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006, 16, 2380-2386.	2.2	71
29	β -Secretase inhibitors: Modification at the P4 position and improvement of inhibitory activity in cultured cells. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006, 16, 4354-4359.	2.2	55
30	A structure-based strategy for discovery of small ligands binding to functionally unknown proteins: Combination of in silico screening and surface plasmon resonance measurements. <i>Proteomics</i> , 2005, 5, 1472-1480.	2.2	48
31	A computational model on the modulation of mitogen-activated protein kinase (MAPK) and Akt pathways in heregulin-induced ErbB signalling. <i>Biochemical Journal</i> , 2003, 373, 451-463.	3.7	220
32	Site-specific incorporation of an unnatural amino acid into proteins in mammalian cells. <i>Nucleic Acids Research</i> , 2002, 30, 4692-4699.	14.5	231
33	Crystal Structure of the Complex of Human Epidermal Growth Factor and Receptor Extracellular Domains. <i>Cell</i> , 2002, 110, 775-787.	28.9	1,013
34	Chimeric receptor analyses of the interactions of the ectodomains of ErbB-1 with epidermal growth factor and of those of ErbB-4 with neuregulin. <i>FEBS Journal</i> , 2002, 269, 2323-2329.	0.2	26
35	The Structure of Physalin T from <i>Physalis alkekengi</i> var. <i>francheti</i> . <i>Journal of Asian Natural Products Research</i> , 2001, 3, 199-205.	1.4	28
36	Further evidence for the involvement of insulin receptor substrates in epidermal growth factor-induced activation of phosphatidylinositol 3-kinase. <i>FEBS Journal</i> , 2001, 268, 4158-4168.	0.2	15

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37	Characterization of the N-Oligosaccharides Attached to the Atypical Asn-X-Cys Sequence of Recombinant Human Epidermal Growth Factor Receptor. <i>Journal of Biochemistry</i> , 2000, 127, 65-72.	1.7	63
38	Phosphodiester bond cleavage mediated by a cyclic β -sheet peptide-based dinuclear zinc(ii) complex. <i>Chemical Communications</i> , 2000, , 1315-1316.	4.1	39
39	Title is missing!. <i>International Journal of Peptide Research and Therapeutics</i> , 1998, 5, 5-12.	0.1	1
40	Stereochemistry of protected ornithine side chains in gramicidin S derivatives and their resistance to N-methylation. <i>International Journal of Peptide Research and Therapeutics</i> , 1998, 5, 5-12.	0.1	2
41	Such Hydrophobic Peptides as Dansylated Mastoparan Can Elevate the Fertilization Membrane of Sea Urchin Eggs. <i>Biochemical and Biophysical Research Communications</i> , 1995, 215, 828-834.	2.1	4
42	General method for preparing altrosides from 2,3-manno-epoxides and its application to synthesis of alternative β -cyclodextrin with an altroside as the constituent of macrocyclic structure. <i>Tetrahedron Letters</i> , 1994, 35, 9577-9580.	1.4	44
43	Determination of the structures of tris(6-O-mesitylenesulfonyl)-.alpha.-cyclodextrin regioisomers by proton NMR analyses of the corresponding 3,6-anhydrocyclodextrin derivatives. <i>Journal of Organic Chemistry</i> , 1993, 58, 2936-2937.	3.2	13
44	Chemotactic peptide from ropalidian wasp as well as the authentic chemotactic tripeptide stimulates two distinct pathways in neutrophils, but the [LYS7] analog does only one of them. <i>Biochemical and Biophysical Research Communications</i> , 1991, 175, 165-172.	2.1	13
45	Preparation, Stereochemistry, and Antibacterial Activity of Gramicidin S Analogs Containing N-Methyl Groups. <i>Bulletin of the Chemical Society of Japan</i> , 1991, 64, 35-41.	3.2	14
46	A 1H-NMR study of the solution conformation of Icaria chemotactic peptide and its [Lys7] analog: Effects on the physiological activity of a substitution of proline to lysine at position 7. <i>Biochemical and Biophysical Research Communications</i> , 1990, 168, 596-603.	2.1	2
47	Role of lysine residue at 7th position of wasp chemotactic peptides. <i>Biochemical and Biophysical Research Communications</i> , 1990, 168, 844-849.	2.1	11
48	Structures of autoxidation products of 2-tert-butyl-4-methoxyphenol in aqueous alkaline solution. <i>Journal of Organic Chemistry</i> , 1989, 54, 4215-4217.	3.2	10
49	A New physalin from <i>Physalis alkekengi</i> : structure of physalin L. <i>Phytochemistry</i> , 1987, 26, 3313-3317.	2.9	41
50	A wasp venom mastoparan-induced polyphosphoinositide breakdown in rat peritoneal mast cells. <i>FEBS Letters</i> , 1985, 188, 363-366.	2.8	73
51	Molecular aggregation and conformational change of wasp venom mastoparan as induced by salt in aqueous solution. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 1984, 802, 157-161.	2.4	18
52	SELECTIVE N-METHYLATION OF PEPTIDE BOND. PREPARATION AND PROPERTIES OF [MeOrn2,2â€², D-MePhe4,4â€²]GRAMICIDIN S. <i>Chemistry Letters</i> , 1984, 13, 1835-1836.	1.3	3
53	Synthesis of a wasp venom tetradecapeptide, mastoparan, with a new cleaving system for 4-methoxy-2,3,6-trimethylbenzenesulfonyl (Mtr) amino-protecting group.. <i>Chemical and Pharmaceutical Bulletin</i> , 1984, 32, 2187-2193.	1.3	11