

Anders Bach

List of Publications by Year in descending order

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51
papers

2,094
citations

257450

24
h-index

233421

45
g-index

53
all docs

53
docs citations

53
times ranked

2721
citing authors

#	ARTICLE	IF	CITATIONS
1	Common Structural Basis for Constitutive Activity of the Ghrelin Receptor Family. <i>Journal of Biological Chemistry</i> , 2004, 279, 53806-53817.	3.4	303
2	A high-affinity, dimeric inhibitor of PSD-95 bivalently interacts with PDZ1-2 and protects against ischemic brain damage. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 3317-3322.	7.1	162
3	A Conserved Aromatic Lock for the Tryptophan Rotameric Switch in TM-VI of Seven-transmembrane Receptors. <i>Journal of Biological Chemistry</i> , 2010, 285, 3973-3985.	3.4	126
4	Nonpeptide and Peptide Growth Hormone Secretagogues Act Both as Ghrelin Receptor Agonist and as Positive or Negative Allosteric Modulators of Ghrelin Signaling. <i>Molecular Endocrinology</i> , 2005, 19, 2400-2411.	3.7	111
5	Identification of a small-molecule inhibitor of the PICK1 PDZ domain that inhibits hippocampal LTP and LTD. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 413-418.	7.1	100
6	Ghrelin Receptor Inverse Agonists: Identification of an Active Peptide Core and Its Interaction Epitopes on the Receptor. <i>Molecular Pharmacology</i> , 2006, 70, 936-946.	2.3	82
7	Cell-Permeable and Plasma-Stable Peptidomimetic Inhibitors of the Postsynaptic Density-95/ <i>N-Methyl-D-Aspartate</i> Receptor Interaction. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 1333-1346.	6.4	81
8	Acid Ceramidase in Melanoma. <i>Journal of Biological Chemistry</i> , 2016, 291, 2422-2434.	3.4	72
9	Non-covalent Small-Molecule Kelch-like ECH-Associated Protein 1 "Nuclear Factor Erythroid 2-Related Factor 2 (Keap1 "Nrf2) Inhibitors and Their Potential for Targeting Central Nervous System Diseases. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 8088-8103.	6.4	71
10	Ligand binding by PDZ domains. <i>BioFactors</i> , 2012, 38, 338-348.	5.4	66
11	A Comparative Assessment Study of Known Small-Molecule Keap1 "Nrf2 Protein "Protein Interaction Inhibitors: Chemical Synthesis, Binding Properties, and Cellular Activity. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 8028-8052.	6.4	66
12	Modified Peptides as Potent Inhibitors of the Postsynaptic Density-95/ <i>N-Methyl-D-Aspartate</i> Receptor Interaction. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 6450-6459.	6.4	61
13	Design and Synthesis of Highly Potent and Plasma-Stable Dimeric Inhibitors of the PSD "NMDA Receptor Interaction. <i>Angewandte Chemie - International Edition</i> , 2009, 48, 9685-9689.	13.8	55
14	A Sequential Binding Mechanism in a PDZ Domain. <i>Biochemistry</i> , 2009, 48, 7089-7097.	2.5	46
15	Unique Interaction Pattern for a Functionally Biased Ghrelin Receptor Agonist. <i>Journal of Biological Chemistry</i> , 2011, 286, 20845-20860.	3.4	42
16	Side-Chain Interactions Form Late and Cooperatively in the Binding Reaction between Disordered Peptides and PDZ Domains. <i>Journal of the American Chemical Society</i> , 2012, 134, 599-605.	13.7	41
17	Energetic Pathway Sampling in a Protein Interaction Domain. <i>Structure</i> , 2013, 21, 1193-1202.	3.3	38
18	The dynamics of linear polyubiquitin. <i>Science Advances</i> , 2020, 6, .	10.3	38

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19	Structure-activity relationships of a small-molecule inhibitor of the PDZ domain of PICK1. <i>Organic and Biomolecular Chemistry</i> , 2010, 8, 4281.	2.8	31
20	GHB analogs confer neuroprotection through specific interaction with the CaMKII α hub domain. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021, 118, .	7.1	31
21	Deconstructing Noncovalent Kelch-like ECH-Associated Protein 1 (Keap1) Inhibitors into Fragments to Reconstruct New Potent Compounds. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 4623-4661.	6.4	30
22	Deciphering the Kinetic Binding Mechanism of Dimeric Ligands Using a Potent Plasma-stable Dimeric Inhibitor of Postsynaptic Density Protein-95 as an Example. <i>Journal of Biological Chemistry</i> , 2010, 285, 28252-28260.	3.4	29
23	Biochemical investigations of the mechanism of action of small molecules ZL006 and IC87201 as potential inhibitors of the nNOS-PDZ/PSD-95-PDZ interactions. <i>Scientific Reports</i> , 2015, 5, 12157.	3.3	29
24	Benzoxazolone Carboxamides as Potent Acid Ceramidase Inhibitors: Synthesis and Structure-Activity Relationship (SAR) Studies. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 9258-9272.	6.4	29
25	PSD-95 uncoupling from NMDA receptors by Tat-N-dimer ameliorates neuronal depolarization in cortical spreading depression. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2017, 37, 1820-1828.	4.3	27
26	Developing Inhibitors of the p47phox-p22phox Protein-Protein Interaction by Fragment-Based Drug Discovery. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 1156-1177.	6.4	25
27	Benzoxazolone Carboxamides: Potent and Systemically Active Inhibitors of Intracellular Acid Ceramidase. <i>Angewandte Chemie - International Edition</i> , 2014, 54, n/a-n/a.	13.8	23
28	Conjugation of Therapeutic PSD-95 Inhibitors to the Cell-Penetrating Peptide Tat Affects Blood-Brain Barrier Adherence, Uptake, and Permeation. <i>Pharmaceutics</i> , 2020, 12, 661.	4.5	22
29	Selectivity, efficacy and toxicity studies of UCCB01-144, a dimeric neuroprotective PSD-95 inhibitor. <i>Neuropharmacology</i> , 2019, 150, 100-111.	4.1	21
30	UCCB01-125, a dimeric inhibitor of PSD-95, reduces inflammatory pain without disrupting cognitive or motor performance: Comparison with the NMDA receptor antagonist MK-801. <i>Neuropharmacology</i> , 2013, 67, 193-200.	4.1	20
31	A high-affinity, bivalent PDZ domain inhibitor complexes PICK1 to alleviate neuropathic pain. <i>EMBO Molecular Medicine</i> , 2020, 12, e11248.	6.9	20
32	Biophysical Characterization of the Complex between Human Papillomavirus E6 Protein and Synapse-associated Protein 97. <i>Journal of Biological Chemistry</i> , 2011, 286, 3597-3606.	3.4	18
33	Targeting Protein-Protein Interactions with Trimeric Ligands: High Affinity Inhibitors of the MAGUK Protein Family. <i>PLoS ONE</i> , 2015, 10, e0117668.	2.5	17
34	Molecular architecture of the Jumonji C family histone demethylase KDM5B. <i>Scientific Reports</i> , 2019, 9, 4019.	3.3	16
35	Rigidified Clicked Dimeric Ligands for Studying the Dynamics of the PDZ1 Supramodule of PSD-95. <i>ChemBioChem</i> , 2015, 16, 64-69.	2.6	15
36	In vitro and in vivo effects of a novel dimeric inhibitor of PSD-95 on excitotoxicity and functional recovery after experimental traumatic brain injury. <i>European Journal of Neuroscience</i> , 2017, 45, 238-248.	2.6	14

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37	A Fluorescence Polarization Based Screening Assay for Identification of Small Molecule Inhibitors of the PICK1 PDZ Domain. <i>Combinatorial Chemistry and High Throughput Screening</i> , 2011, 14, 590-600.	1.1	12
38	PDZ Domain-Mediated Interactions of G Protein-Coupled Receptors with Postsynaptic Density Protein 95: Quantitative Characterization of Interactions. <i>PLoS ONE</i> , 2013, 8, e63352.	2.5	11
39	Design, Synthesis, and Characterization of Fatty Acid Derivatives of a Dimeric Peptide-Based Postsynaptic Density-95 (PSD-95) Inhibitor. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 1575-1580.	6.4	10
40	Selective release of gastrointestinal hormones induced by an orally active GPR39 agonist. <i>Molecular Metabolism</i> , 2021, 49, 101207.	6.5	9
41	Design and synthesis of triazole-based peptidomimetics of a PSD-95 PDZ domain inhibitor. <i>MedChemComm</i> , 2016, 7, 531-536.	3.4	8
42	Targeting Oxidative Stress in Stroke. <i>Springer Series in Translational Stroke Research</i> , 2017, , 203-250.	0.1	8
43	Detecting Protein-Protein Interactions in Living Cells: Development of a Bioluminescence Resonance Energy Transfer Assay to Evaluate the PSD-95/NMDA Receptor Interaction. <i>Neurochemical Research</i> , 2009, 34, 1729-1737.	3.3	7
44	The European Federation for Medicinal Chemistry and Chemical Biology (EFMC) Best Practice Initiative: Phenotypic Drug Discovery. <i>ChemMedChem</i> , 2021, 16, 1737-1740.	3.2	7
45	Effects of the dimeric PSD-95 inhibitor UCCB01-144 in mouse models of pain, cognition and motor function. <i>European Journal of Pharmacology</i> , 2016, 780, 166-173.	3.5	6
46	Interaction partners of PSD-93 studied by X-ray crystallography and fluorescence polarization spectroscopy. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2013, 69, 587-594.	2.5	5
47	Effects of Dimeric PSD-95 Inhibition on Excitotoxic Cell Death and Outcome After Controlled Cortical Impact in Rats. <i>Neurochemical Research</i> , 2017, 42, 3401-3413.	3.3	5
48	Identification and Optimization of Novel Small-Molecule Cas9 Inhibitors by Cell-Based High-Throughput Screening. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 3266-3305.	6.4	5
49	Expedient Synthesis of 1,3-Substituted Benzene Peptidomimetics. <i>Synthesis</i> , 2011, 2011, 807-815.	2.3	2
50	Effects of the dimeric PSD-95 inhibitor UCCB01-144 on functional recovery after fimbria-fornix transection in rats. <i>Pharmacology Biochemistry and Behavior</i> , 2017, 161, 62-67.	2.9	2
51	Identification of Novel Fragments Binding to the PDZ1 Domain of PSD-95. <i>ChemMedChem</i> , 2021, 16, 949-954.	3.2	1