Kasper B Hansen

List of Publications by Year
in descending order

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80 papers	7,011 citations	126907 33 h-index	74163 75 g-index
82 all docs	82 docs citations	82 times ranked	7120 citing authors

#	Article	IF	Citations
1	Derivatives of (R)-3-(5-Furanyl)carboxamido-2-aminopropanoic Acid as Potent NMDA Receptor Glycine Site Agonists with GluN2 Subunit-Specific Activity. Journal of Medicinal Chemistry, 2022, 65, 734-746.	6.4	5
2	Synaptic Dysfunction by Mutations in GRIN2B: Influence of Triheteromeric NMDA Receptors on Gain-of-Function and Loss-of-Function Mutant Classification. Brain Sciences, 2022, 12, 789.	2.3	9
3	Stereoselective synthesis of novel 2â€²-(S)-CCG-IV analogues as potent NMDA receptor agonists. European Journal of Medicinal Chemistry, 2021, 212, 113099.	5.5	3
4	Distinct GluN1 and GluN2 Structural Determinants for Subunit-Selective Positive Allosteric Modulation of N-Methyl-<scp>d</scp>-aspartate Receptors. ACS Chemical Neuroscience, 2021, 12, 79-98.	3.5	10
5	Structure, Function, and Pharmacology of Glutamate Receptor Ion Channels. Pharmacological Reviews, 2021, 73, 1469-1658.	16.0	237
6	Negative allosteric modulation of GluN1/GluN3 NMDA receptors. Neuropharmacology, 2020, 176, 108117.	4.1	17
7	Utilizing a C(sp3)â€“H Activation Strategy and Structureâ€“Activity Relationship Studies at the Ionotropic Glutamate Receptors. ACS Chemical Neuroscience, 2020, 11, 674-701.	3.5	8
8	Improved synthetic route for the GluN2-specific NMDA receptor glycine site agonist AICP. Tetrahedron Letters, 2020, 61, 151653.	1.4	4
9	PTC-174, a positive allosteric modulator of NMDA receptors containing GluN2C or GluN2D subunits. Neuropharmacology, 2020, 173, 107971.	4.1	13
10	Functional and pharmacological properties of triheteromeric GluN1/2B/2D NMDA receptors. Journal of Physiology, 2019, 597, 5495-5514.	2.9	38
11	Design and Synthesis of 2,3-trans-Proline Analogues as Ligands for Ionotropic Glutamate Receptors and Excitatory Amino Acid Transporters. ACS Chemical Neuroscience, 2019, 10, 2989-3007.	3.5	4
12	Use of the 4-Hydroxytriazole Moiety as a Bioisosteric Tool in the Development of Ionotropic Glutamate Receptor Ligands. Journal of Medicinal Chemistry, 2019, 62, 4467-4482.	6.4	18
13	Functional assessment of triheteromeric NMDA receptors containing a human variant associated with epilepsy. Journal of Physiology, 2019, 597, 1691-1704.	2.9	20
14	Properties of TriheteromericN-Methyl-d-Aspartate Receptors Containing Two Distinct GluN1 Isoforms. Molecular Pharmacology, 2018, 93, 453-467.	2.3	24
15	Structure, function, and allosteric modulation of NMDA receptors. Journal of General Physiology, 2018, 150, 1081-1105.	1.9	363
16	Triheteromeric GluN1/GluN2A/GluN2C NMDARs with Unique Single-Channel Properties Are the Dominant Receptor Population in Cerebellar Granule Cells. Neuron, 2018, 99, 315-328.e5.	8.1	42
17	Subtype-Specific Agonists for NMDA Receptor Glycine Binding Sites. ACS Chemical Neuroscience, 2017, 8, 1681-1687.	3.5	19
18	Identification of AICP as a GluN2C-Selective N-Methyl-d-Aspartate Receptor Superagonist at the GluN1 Glycine Site. Molecular Pharmacology, 2017, 92, 151-161.	2.3	16

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19	Allosteric Interactions between NMDA Receptor Subunits Shape the Developmental Shift in Channel Properties. Neuron, 2017, 94, 58-64.e3.	8.1	38
20	NMDA Receptors in the Central Nervous System. Methods in Molecular Biology, 2017, 1677, 1-80.	0.9	105
21	Selective Cell-Surface Expression of Triheteromeric NMDA Receptors. Methods in Molecular Biology, 2017, 1677, 145-162.	0.9	11
22	Structural basis of subunit selectivity for competitive NMDA receptor antagonists with preference for GluN2A over GluN2B subunits. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E6942-E6951.	7.1	33
23	Augmentation of Anticancer Drug Efficacy in Murine Hepatocellular Carcinoma Cells by a Peripherally Acting Competitive N-Methyl-<scp>d</scp>-aspartate (NMDA) Receptor Antagonist. Journal of Medicinal Chemistry, 2017, 60, 9885-9904.	6.4	27
24	Structural Basis for Negative Allosteric Modulation of GluN2A-Containing NMDA Receptors. Neuron, 2016, 91, 1316-1329.	8.1	74
25	Development of Radiolabeled Ligands Targeting the Glutamate Binding Site of the N-Methyl-<scp>d</scp>-aspartate Receptor as Potential Imaging Agents for Brain. Journal of Medicinal Chemistry, 2016, 59, 11110-11119.	6.4	16
26	Pharmacology of triheteromeric N-Methyl-d-Aspartate Receptors. Neuroscience Letters, 2016, 617, 240-246.	2.1	24
27	Pharmacology and Structural Analysis of Ligand Binding to the Orthosteric Site of Glutamate-Like GluD2 Receptors. Molecular Pharmacology, 2016, 89, 253-262.	2.3	26
28	Binding of ArgTX-636 in the NMDA Receptor Ion Channel. Journal of Molecular Biology, 2015, 427, 176-189.	4.2	13
29	Structural Determinants and Mechanism of Action of a GluN2C-selective NMDA Receptor Positive Allosteric Modulator. Molecular Pharmacology, 2014, 86, 548-560.	2.3	69
30	Functional analysis of a de novo GRIN2A missense mutation associated with early-onset epileptic encephalopathy. Nature Communications, 2014, 5, 3251.	12.8	128
31	Distinct Functional and Pharmacological Properties of Triheteromeric GluN1/GluN2A/GluN2B NMDA Receptors. Neuron, 2014, 81, 1084-1096.	8.1	246
32	Design, Synthesis, and Structureâ€“Activity Relationship of a Novel Series of GluN2C-Selective Potentiators. Journal of Medicinal Chemistry, 2014, 57, 2334-2356.	6.4	43
33	Mechanistic twists and turns. Nature Chemical Biology, 2014, 10, 698-699.	8.0	2
34	A Human Mutation in the M4 Helix of GluN2A Accelerates Forward Gating Transitions in NMDA Receptors. Biophysical Journal, 2014, 106, 150a.	0.5	0
35	Crystal Structure and Pharmacological Characterization of a Novel N-Methyl-d-aspartate (NMDA) Receptor Antagonist at the GluN1 Glycine Binding Site. Journal of Biological Chemistry, 2013, 288, 33124-33135.	3.4	22
36	Modal gating of GluN1/GluN2D NMDA receptors. Neuropharmacology, 2013, 71, 184-190.	4.1	15

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37	Structure-based discovery of antagonists for GluN3-containing N-methyl-d-aspartate receptors. Neuropharmacology, 2013, 75, 324-336.	4.1	36
38	Development of 2â€²-Substituted (2S,1â€²R,2â€²S)-2-(Carboxycyclopropyl)glycine Analogues as PotentN-Methyl-d-aspartic Acid Receptor Agonists. Journal of Medicinal Chemistry, 2013, 56, 4071-4081.	6.4	9
39	Structural Determinants of Agonist Efficacy at the Glutamate Binding Site ofN-Methyl-d-Aspartate Receptors. Molecular Pharmacology, 2013, 84, 114-127.	2.3	76
40	Subunit-Selective Allosteric Inhibition of Glycine Binding to NMDA Receptors. Journal of Neuroscience, 2012, 32, 6197-6208.	3.6	99
41	Molecular pharmacology of human NMDA receptors. Neurochemistry International, 2012, 61, 601-609.	3.8	74
42	GluN1 splice variant control of GluN1/GluN2D NMDA receptors. Journal of Physiology, 2012, 590, 3857-3875.	2.9	52
43	6.2 Structure-Function Correlates of Glutamate-Gated Ion Channels. , 2012, , 4-30.		0
44	The use ofXenopusoocytes in drug screening. Expert Opinion on Drug Discovery, 2011, 6, 141-153.	5.0	23
45	How Glutamate Receptor Subunits Mix and Match: Details Uncovered. Neuron, 2011, 71, 198-200.	8.1	2
46	Structural and Mechanistic Determinants of a Novel Site for Noncompetitive Inhibition of GluN2D-Containing NMDA Receptors. Journal of Neuroscience, 2011, 31, 3650-3661.	3.6	67
47	Mechanism for Noncompetitive Inhibition by Novel GluN2C/D N-Methyl-d-aspartate Receptor Subunit-Selective Modulators. Molecular Pharmacology, 2011, 80, 782-795.	2.3	89
48	Quinazolin-4-one Derivatives: A Novel Class of Noncompetitive NR2C/D Subunit-Selective N-Methyl-<scp>d</scp>-aspartate Receptor Antagonists. Journal of Medicinal Chemistry, 2010, 53, 5476-5490.	6.4	83
49	Novel 3â€Carboxyâ€•and 3â€Phosphonopyrazoline Amino Acids as Potent and Selective NMDA Receptor Antagonists: Design, Synthesis, and Pharmacological Characterization. ChemMedChem, 2010, 5, 1465-1475.	3.2	22
50	Partial Agonists and Subunit Selectivity at NMDA Receptors. Chemistry - A European Journal, 2010, 16, 13910-13918.	3.3	8
51	Mutational Mapping and Modeling of the Binding Site for (S)-Citalopram in the Human Serotonin Transporter. Journal of Biological Chemistry, 2010, 285, 2051-2063.	3.4	91
52	Implementation of a Fluorescence-Based Screening Assay Identifies Histamine H3 Receptor Antagonists Clobenpropit and Iodophenpropit as Subunit-Selective N-Methyl-d-Aspartate Receptor Antagonists. Journal of Pharmacology and Experimental Therapeutics, 2010, 333, 650-662.	2.5	40
53	A subunit-selective potentiator of NR2C- and NR2D-containing NMDA receptors. Nature Communications, 2010, 1, 90.	12.8	137
54	Control of Assembly and Function of Glutamate Receptors by the Amino-Terminal Domain. Molecular Pharmacology, 2010, 78, 535-549.	2.3	95

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55	Subunit-Specific Activation of NMDA Receptors. Biophysical Journal, 2010, 98, 525a-526a.	0.5	2
56	4-Hydroxy-1,2,5-oxadiazol-3-yl Moiety as Bioisoster of the Carboxy Function. Synthesis, Ionization Constants, and Molecular Pharmacological Characterization at Ionotropic Glutamate Receptors of Compounds Related to Glutamate and Its Homologues. Journal of Medicinal Chemistry, 2010, 53, 4110-4118.	6.4	24
57	Glutamate Receptor Ion Channels: Structure, Regulation, and Function. Pharmacological Reviews, 2010, 62, 405-496.	16.0	2,973
58	Activation of Recombinant Rat GluN1/gluN2D NMDA Receptors. Biophysical Journal, 2010, 98, 525a.	0.5	1
59	Modulation of the Dimer Interface at Ionotropic Glutamate-Like Receptor Î´2 by d-Serine and Extracellular Calcium. Journal of Neuroscience, 2009, 29, 907-917.	3.6	57
60	Location of the Antidepressant Binding Site in the Serotonin Transporter. Journal of Biological Chemistry, 2009, 284, 10276-10284.	3.4	105
61	Stereocontrolled Synthesis and Pharmacological Evaluation of Azetidineâ€2,3â€Dicarboxylic Acids at NMDA Receptors. ChemMedChem, 2009, 4, 110-117.	3.2	10
62	FLIPRÂ® Assays of Intracellular Calcium in GPCR Drug Discovery. Methods in Molecular Biology, 2009, 552, 269-278.	0.9	23
63	An allosteric binding site at the human serotonin transporter mediates the inhibition of escitalopram by R-citalopram: Kinetic binding studies with the ALI/VFLâ€“SI/TT mutant. Neuroscience Letters, 2009, 462, 207-212.	2.1	41
64	Control of NMDA Receptor Function by the NR2 Subunit Amino-Terminal Domain. Journal of Neuroscience, 2009, 29, 12045-12058.	3.6	189
65	Xenopus Oocyte Electrophysiology in GPCR Drug Discovery. Methods in Molecular Biology, 2009, 552, 343-357.	0.9	18
66	Structural Correlates of Ionotropic Glutamate Receptor Function. , 2008, , 247-297.		0
67	N-Hydroxypyrazolyl Glycine Derivatives as Selective N-Methyl-d-aspartic Acid Receptor Ligands. Journal of Medicinal Chemistry, 2008, 51, 4179-4187.	6.4	19
68	Pharmacological Characterization of Ligands at Recombinant NMDA Receptor Subtypes by Electrophysiological Recordings and Intracellular Calcium Measurements. Combinatorial Chemistry and High Throughput Screening, 2008, 11, 304-315.	1.1	31
69	Structure and Function of the NMDA Receptor. , 2008, , 289-316.		11
70	Ionotropic glutamate-like receptor Î´2 binds <scp>d</scp> -serine and glycine. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 14116-14121.	7.1	138
71	Pharmacological characterization of mouse GPRC6A, an L -Î± -amino-acid receptor modulated by divalent cations. British Journal of Pharmacology, 2007, 150, 798-807.	5.4	74
72	Structural aspects of AMPA receptor activation, desensitization and deactivation. Current Opinion in Neurobiology, 2007, 17, 281-288.	4.2	68

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73	Subunit-Specific Agonist Activity at NR2A-, NR2B-, NR2C-, and NR2D-ContainingN-Methyl-d-aspartate Glutamate Receptors. Molecular Pharmacology, 2007, 72, 907-920.	2.3	151
74	Synthesis and pharmacology of glutamate receptor ligands: new isothiazole analogues of ibotenic acid. Organic and Biomolecular Chemistry, 2007, 5, 463-471.	2.8	21
75	Deorphanization of GPRC6A: A Promiscuous l-Î±-Amino Acid Receptor with Preference for Basic Amino Acids. Molecular Pharmacology, 2005, 67, 589-597.	2.3	194
76	Tweaking Agonist Efficacy at N-Methyl-d-aspartate Receptors by Site-Directed Mutagenesis. Molecular Pharmacology, 2005, 68, 1510-1523.	2.3	24
77	Synthesis, Binding Affinity at Glutamic Acid Receptors, Neuroprotective Effects, and Molecular Modeling Investigation of Novel Dihydroisoxazole Amino Acids. Journal of Medicinal Chemistry, 2005, 48, 6315-6325.	6.4	43
78	The respective N-hydroxypyrazole analogues of the classical glutamate receptor ligands ibotenic acid and (RS)-2-amino-2-(3-hydroxy-5-methyl-4-isoxazolyl)acetic acid. European Journal of Pharmacology, 2004, 499, 35-44.	3.5	23
79	(S)-2-Amino-3-(3-hydroxy-7,8-dihydro-6H-cyclohepta[d]isoxazol-4-yl)propionic Acid, a Potent and Selective Agonist at the GluR5 Subtype of Ionotropic Glutamate Receptors. Synthesis, Modeling, and Molecular Pharmacology. Journal of Medicinal Chemistry, 2003, 46, 1350-1358.	6.4	26
80	Activation of NMDA receptors and postsynaptic events. Reactome - A Curated Knowledgebase of Biological Pathways, 0, 67, .	0.0	0