

Kristoffer Sahlholm

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/734954/publications.pdf>

Version: 2024-02-01

43
papers

860
citations

471509
17
h-index

501196
28
g-index

44
all docs

44
docs citations

44
times ranked

1181
citing authors

#	ARTICLE	IF	CITATIONS
1	Facilitated Anion Transport Induces Hyperpolarization of the Cell Membrane That Triggers Differentiation and Cell Death in Cancer Stem Cells. <i>Journal of the American Chemical Society</i> , 2015, 137, 15892-15898.	13.7	109
2	The dopamine stabilizers ACR16 and (âˆ²)-OSU6162 display nanomolar affinities at the Î¶f-1 receptor. <i>Molecular Psychiatry</i> , 2013, 18, 12-14.	7.9	70
3	Pridopidine selectively occupies sigma-1 rather than dopamine D2 receptors at behaviorally active doses. <i>Psychopharmacology</i> , 2015, 232, 3443-3453.	3.1	55
4	Evidence for oligomerization between GABA_B receptors and GIRK channels containing the GIRK1 and GIRK3 subunits. <i>European Journal of Neuroscience</i> , 2010, 32, 1265-1277.	2.6	52
5	The fast-off hypothesis revisited: A functional kinetic study of antipsychotic antagonism of the dopamine D2 receptor. <i>European Neuropsychopharmacology</i> , 2016, 26, 467-476.	0.7	38
6	A Neanderthal Sodium Channel Increases Pain Sensitivity in Present-Day Humans. <i>Current Biology</i> , 2020, 30, 3465-3469.e4.	3.9	33
7	Agonist-specific voltage sensitivity at the dopamine D2S receptor â€“ Molecular determinants and relevance to therapeutic ligands. <i>Neuropharmacology</i> , 2011, 61, 937-949.	4.1	31
8	Voltageâ€dependence of the human dopamine D₂ receptor. <i>Synapse</i> , 2008, 62, 476-480.	1.2	29
9	Antipsychotic-Like Efficacy of Dopamine D2 Receptor-Biased Ligands is Dependent on Adenosine A2A Receptor Expression. <i>Molecular Neurobiology</i> , 2018, 55, 4952-4958.	4.0	28
10	Highly Selective Dopamine D₃ Receptor Antagonists with Arylated Diazaspiro Alkane Cores. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 9905-9910.	6.4	27
11	Behavioral control by striatal adenosine A_{2A}â€dopamine D₂ receptor heteromers. <i>Genes, Brain and Behavior</i> , 2018, 17, e12432.	2.2	27
12	A truncated Kv1.1 protein in the brain of the megencephaly mouse: expression and interaction. <i>BMC Neuroscience</i> , 2005, 6, 65.	1.9	25
13	Cocaine produces D2R-mediated conformational changes in the adenosine A2AR-dopamine D2R heteromer. <i>Biochemical and Biophysical Research Communications</i> , 2010, 394, 988-992.	2.1	25
14	Decreased striatal adenosine A2A-dopamine D2 receptor heteromerization in schizophrenia. <i>Neuropsychopharmacology</i> , 2021, 46, 665-672.	5.4	24
15	Voltage-sensitivity at the human dopamine D2S receptor is agonist-specific. <i>Biochemical and Biophysical Research Communications</i> , 2008, 377, 1216-1221.	2.1	23
16	Differential voltage-sensitivity of D2-like dopamine receptors. <i>Biochemical and Biophysical Research Communications</i> , 2008, 374, 496-501.	2.1	20
17	Voltage sensitivities and deactivation kinetics of histamine H3 and H4 receptors. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2012, 1818, 3081-3089.	2.6	19
18	G Protein-Gated Inwardly Rectifying Potassium Channel Subunits 1 and 2 are Down-Regulated in Rat Dorsal Root Ganglion Neurons and Spinal Cord after Peripheral Axotomy. <i>Molecular Pain</i> , 2015, 11, s12990-015-0044.	2.1	18

#	ARTICLE	IF	CITATIONS
19	Typical and atypical antipsychotics do not differ markedly in their reversibility of antagonism of the dopamine D2 receptor. <i>International Journal of Neuropsychopharmacology</i> , 2014, 17, 149-155.	2.1	16
20	Sigma-2 receptor binding is decreased in female, but not male, APP/PS1 mice. <i>Biochemical and Biophysical Research Communications</i> , 2015, 460, 439-445.	2.1	16
21	The Beta-Arrestin-Biased Dopamine D2 Receptor Ligand, UNC9994, Is a Partial Agonist at G-Protein-Mediated Potassium Channel Activation. <i>International Journal of Neuropsychopharmacology</i> , 2018, 21, 1102-1108.	2.1	15
22	Leveraging a Low-Affinity Diazaspiro Orthosteric Fragment to Reduce Dopamine D ₃ Receptor (D ₃ R) Ligand Promiscuity across Highly Conserved Aminergic G-Protein-Coupled Receptors (GPCRs). <i>Journal of Medicinal Chemistry</i> , 2019, 62, 5132-5147.	6.4	15
23	Remote local photoactivation of morphine produces analgesia without opioid-related adverse effects. <i>British Journal of Pharmacology</i> , 2023, 180, 958-974.	5.4	15
24	Voltage-Dependent Dopamine Potency at D1-Like Dopamine Receptors. <i>Frontiers in Pharmacology</i> , 2020, 11, 581151.	3.5	12
25	The human histamine H3 receptor couples to GIRK channels in <i>Xenopus</i> oocytes. <i>European Journal of Pharmacology</i> , 2007, 567, 206-210.	3.5	11
26	Point mutation of a conserved aspartate, D69, in the muscarinic M ₂ receptor does not modify voltage-sensitive agonist potency. <i>Biochemical and Biophysical Research Communications</i> , 2018, 496, 101-104.	2.1	10
27	Inhibition of Tryptophan Hydroxylases and Monoamine Oxidase-A by the Proton Pump Inhibitor, Omeprazole—In Vitro and In Vivo Investigations. <i>Frontiers in Pharmacology</i> , 2020, 11, 593416.	3.5	10
28	Discovery and biological characterization of a novel scaffold for potent inhibitors of peripheral serotonin synthesis. <i>Future Medicinal Chemistry</i> , 2020, 12, 1461-1474.	2.3	10
29	V374A KCND3 Pathogenic Variant Associated With Paroxysmal Ataxia Exacerbations. <i>Neurology: Genetics</i> , 2021, 7, e546.	1.9	10
30	Pridopidine Reverses Phencyclidine-Induced Memory Impairment. <i>Frontiers in Pharmacology</i> , 2018, 9, 338.	3.5	9
31	The role of RGS protein in agonist-dependent relaxation of GIRK currents in <i>Xenopus</i> oocytes. <i>Biochemical and Biophysical Research Communications</i> , 2011, 415, 509-514.	2.1	8
32	Ligand with Two Modes of Interaction with the Dopamine D ₂ Receptor—An Induced-Fit Mechanism of Insurmountable Antagonism. <i>ACS Chemical Neuroscience</i> , 2020, 11, 3130-3143.	3.5	8
33	Mechanistic insights into dopaminergic and serotonergic neurotransmission—concerted interactions with helices 5 and 6 drive the functional outcome. <i>Chemical Science</i> , 2021, 12, 10990-11003.	7.4	7
34	Antipsychotics with similar association kinetics at dopamine D2 receptors differ in extrapyramidal side-effects. <i>Nature Communications</i> , 2018, 9, 3577.	12.8	6
35	Interaction of Ligands for PET with the Dopamine D3 Receptor: In Silico and In Vitro Methods. <i>Biomolecules</i> , 2021, 11, 529.	4.0	6
36	Dopamine D2 Receptor Agonist Binding Kinetics—Role of a Conserved Serine Residue. <i>International Journal of Molecular Sciences</i> , 2021, 22, 4078.	4.1	5

#	ARTICLE	IF	CITATIONS
37	The role of beta-arrestin2 in shaping fMRI BOLD responses to dopaminergic stimulation. <i>Psychopharmacology</i> , 2017, 234, 2019-2030.	3.1	4
38	Electrophysiology-based analysis of human histamine H4 receptor pharmacology using GIRK channel coupling in <i>Xenopus</i> oocytes. <i>European Journal of Pharmacology</i> , 2008, 591, 52-58.	3.5	3
39	Effects of the Dopamine Stabilizer, Pridopidine, on Basal and Phencyclidine-Induced Locomotion: Role of Dopamine D2 and Sigma-1 Receptors. <i>CNS and Neurological Disorders - Drug Targets</i> , 2018, 17, 522-527.	1.4	3
40	G protein-coupled receptor kinase-2 confers isoform-specific calcium sensitivity to dopamine D ₂ receptor desensitization. <i>FASEB Journal</i> , 2021, 35, e22013.	0.5	3
41	Dopamine receptor heteromers: biasing antipsychotics. <i>Future Medicinal Chemistry</i> , 2018, 10, 2675-2677.	2.3	2
42	Synthetic corticosteroids as tryptophan hydroxylase stabilizers. <i>Future Medicinal Chemistry</i> , 2021, 13, 1465-1474.	2.3	2
43	Evidence for Two Modes of Binding of the Negative Allosteric Modulator SB269,652 to the Dopamine D2 Receptor. <i>Biomedicines</i> , 2022, 10, 22.	3.2	1