Frederik J R Rombouts

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8131793/publications.pdf

Version: 2024-02-01

27	737	567281	526287
papers	citations	15 h-index	27 g-index

31	31	31	1080
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Tau Positron Emission Tomography (PET) Imaging: Past, Present, and Future. Journal of Medicinal Chemistry, 2015, 58, 4365-4382.	6.4	88
2	Preclinical Evaluation of¹⁸F-JNJ64349311, a Novel PET Tracer for Tau Imaging. Journal of Nuclear Medicine, 2017, 58, 975-981.	5.0	72
3	1,4-Oxazine Î²-Secretase 1 (BACE1) Inhibitors: From Hit Generation to Orally Bioavailable Brain Penetrant Leads. Journal of Medicinal Chemistry, 2015, 58, 8216-8235.	6.4	67
4	Benzazaborinines as Novel Bioisosteric Replacements of Naphthalene: Propranolol as an Example. Journal of Medicinal Chemistry, 2015, 58, 9287-9295.	6.4	62
5	New evolutions in the BACE1 inhibitor field from 2014 to 2018. Bioorganic and Medicinal Chemistry Letters, 2019, 29, 761-777.	2.2	57
6	Pyrido[4,3-e][1,2,4]triazolo[4,3-a]pyrazines as Selective, Brain Penetrant Phosphodiesterase 2 (PDE2) Inhibitors. ACS Medicinal Chemistry Letters, 2015, 6, 282-286.	2.8	49
7	Rational design and synthesis of aminopiperazinones as Î²-secretase (BACE) inhibitors. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 7255-7260.	2.2	44
8	Structure-Based Design of a Potent, Selective, and Brain Penetrating PDE2 Inhibitor with Demonstrated Target Engagement. ACS Medicinal Chemistry Letters, 2014, 5, 1049-1053.	2.8	41
9	Discovery of N-(Pyridin-4-yl)-1,5-naphthyridin-2-amines as Potential Tau Pathology PET Tracers for Alzheimerâ€™s Disease. Journal of Medicinal Chemistry, 2017, 60, 1272-1291.	6.4	31
10	Microwave-assisted N-debenzylation of amides with triflic acid. Tetrahedron Letters, 2010, 51, 4815-4818.	1.4	27
11	Discovery of N-(4-[¹⁸F]Fluoro-5-methylpyridin-2-yl)isoquinolin-6-amine (JNJ-64326067), a New Promising Tau Positron Emission Tomography Imaging Tracer. Journal of Medicinal Chemistry, 2019, 62, 2974-2987.	6.4	24
12	Towards selective phosphodiesterase 2A (PDE2A) inhibitors: a patent review (2010 - present). Expert Opinion on Therapeutic Patents, 2016, 26, 933-946.	5.0	22
13	Small-molecule BACE1 inhibitors: a patent literature review (2011 to 2020). Expert Opinion on Therapeutic Patents, 2021, 31, 25-52.	5.0	22
14	Evaluation of a Series of Î²-Secretase 1 Inhibitors Containing Novel Heteroaryl-Fused-Piperazine Amidine Warheads. ACS Medicinal Chemistry Letters, 2019, 10, 1159-1165.	2.8	20
15	Synthesis of 2,1â€Borazaroquinolines and 2,1â€Borazaroisoquinolines from VinylÂaminopyridines and Potassium Organotrifluoroborates by Microwaveâ€Assisted Heating. European Journal of Organic Chemistry, 2015, 2015, 5221-5229.	2.4	17
16	Clinical evaluation of [18F] JNJ-64326067, a novel candidate PET tracer for the detection of tau pathology in Alzheimerâ€™s disease. European Journal of Nuclear Medicine and Molecular Imaging, 2020, 47, 3176-3185.	6.4	17
17	Optimization of 1,4-Oxazine Î²-Secretase 1 (BACE1) Inhibitors Toward a Clinical Candidate. Journal of Medicinal Chemistry, 2018, 61, 5292-5303.	6.4	15
18	Fragment Binding to Î²-Secretase 1 without Catalytic Aspartate Interactions Identified via Orthogonal Screening Approaches. ACS Omega, 2017, 2, 685-697.	3.5	14

#	Article	IF	Citations
19	Structure-Based Approaches to Improving Selectivity through Utilizing Explicit Water Molecules: Discovery of Selective Î²-Secretase (BACE1) Inhibitors over BACE2. Journal of Medicinal Chemistry, 2021, 64, 3075-3085.	6.4	11
20	JNJ-67569762, A 2-Aminotetrahydropyridine-Based Selective BACE1 Inhibitor Targeting the S3 Pocket: From Discovery to Clinical Candidate. Journal of Medicinal Chemistry, 2021, 64, 14175-14191.	6.4	10
21	Regioselective Preparation of 3â€Alkoxyâ€4,5â€difluoroanilines by S_NAr. European Journal of Organic Chemistry, 2012, 2012, 7048-7052.	2.4	7
22	Industrial medicinal chemistry insights: neuroscience hit generation at Janssen. Drug Discovery Today, 2017, 22, 1478-1488.	6.4	5
23	Discovery of Extremely Selective Fused Pyridine-Derived Î²-Site Amyloid Precursor Protein-Cleaving Enzyme (BACE1) Inhibitors with High In Vivo Efficacy through 10s Loop Interactions. Journal of Medicinal Chemistry, 2021, 64, 14165-14174.	6.4	4
24	Expedient Access to Cyanated Nâ€Heterocycles by Direct Flowâ€Electrochemical C(sp²)â^'H Activation. Chemistry - A European Journal, 2022, 28, .	3.3	4
25	3,3-Difluoro-3,4,5,6-tetrahydropyridin-2-amines: Potent and permeable BACE-1 inhibitors. Bioorganic and Medicinal Chemistry Letters, 2020, 30, 126999.	2.2	3
26	A Brain-Penetrant and Bioavailable Pyrazolopiperazine BACE1 Inhibitor Elicits Sustained Reduction of Amyloid Î² In Vivo. ACS Medicinal Chemistry Letters, 2022, 13, 76-83.	2.8	3
27	Modulating physicochemical properties of tetrahydropyridine-2-amine BACE1 inhibitors with electron-withdrawing groups: A systematic study. European Journal of Medicinal Chemistry, 2022, 228, 114028.	5.5	0