

Frederik J R Rombouts

List of Publications by Year in descending order

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27
papers

737
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567281

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31
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31
times ranked

1080
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Expedient Access to Cyanated N-Heterocycles by Direct Flow Electrochemical C(sp ²) ^H Activation. <i>Chemistry - A European Journal</i> , 2022, 28, . | 3.3 | 4 |
| 2 | Modulating physicochemical properties of tetrahydropyridine-2-amine BACE1 inhibitors with electron-withdrawing groups: A systematic study. <i>European Journal of Medicinal Chemistry</i> , 2022, 228, 114028. | 5.5 | 0 |
| 3 | A Brain-Penetrant and Bioavailable Pyrazolopiperazine BACE1 Inhibitor Elicits Sustained Reduction of Amyloid β In Vivo. <i>ACS Medicinal Chemistry Letters</i> , 2022, 13, 76-83. | 2.8 | 3 |
| 4 | Small-molecule BACE1 inhibitors: a patent literature review (2011 to 2020). <i>Expert Opinion on Therapeutic Patents</i> , 2021, 31, 25-52. | 5.0 | 22 |
| 5 | Structure-Based Approaches to Improving Selectivity through Utilizing Explicit Water Molecules: Discovery of Selective β -Secretase (BACE1) Inhibitors over BACE2. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 3075-3085. | 6.4 | 11 |
| 6 | JNJ-67569762, A 2-Aminotetrahydropyridine-Based Selective BACE1 Inhibitor Targeting the S3 Pocket: From Discovery to Clinical Candidate. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 14175-14191. | 6.4 | 10 |
| 7 | Discovery of Extremely Selective Fused Pyridine-Derived β -Site Amyloid Precursor Protein-Cleaving Enzyme (BACE1) Inhibitors with High In Vivo Efficacy through 10s Loop Interactions. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 14165-14174. | 6.4 | 4 |
| 8 | Clinical evaluation of [¹⁸ F] JNJ-64326067, a novel candidate PET tracer for the detection of tau pathology in Alzheimer's disease. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2020, 47, 3176-3185. | 6.4 | 17 |
| 9 | 3,3-Difluoro-3,4,5,6-tetrahydropyridin-2-amines: Potent and permeable BACE-1 inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 126999. | 2.2 | 3 |
| 10 | Evaluation of a Series of β -Secretase 1 Inhibitors Containing Novel Heteroaryl-Fused-Piperazine Amidine Warheads. <i>ACS Medicinal Chemistry Letters</i> , 2019, 10, 1159-1165. | 2.8 | 20 |
| 11 | Discovery of <i>N</i> -(4-[(¹⁸ F)Fluoro-5-methylpyridin-2-yl]isoquinolin-6-amine (JNJ-64326067), a New Promising Tau Positron Emission Tomography Imaging Tracer. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 2974-2987. | 6.4 | 24 |
| 12 | New evolutions in the BACE1 inhibitor field from 2014 to 2018. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 761-777. | 2.2 | 57 |
| 13 | Optimization of 1,4-Oxazine β -Secretase 1 (BACE1) Inhibitors Toward a Clinical Candidate. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 5292-5303. | 6.4 | 15 |
| 14 | Discovery of <i>N</i> -(Pyridin-4-yl)-1,5-naphthyridin-2-amines as Potential Tau Pathology PET Tracers for Alzheimer's Disease. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 1272-1291. | 6.4 | 31 |
| 15 | Preclinical Evaluation of [¹⁸ F]JNJ64349311, a Novel PET Tracer for Tau Imaging. <i>Journal of Nuclear Medicine</i> , 2017, 58, 975-981. | 5.0 | 72 |
| 16 | Fragment Binding to β -Secretase 1 without Catalytic Aspartate Interactions Identified via Orthogonal Screening Approaches. <i>ACS Omega</i> , 2017, 2, 685-697. | 3.5 | 14 |
| 17 | Industrial medicinal chemistry insights: neuroscience hit generation at Janssen. <i>Drug Discovery Today</i> , 2017, 22, 1478-1488. | 6.4 | 5 |
| 18 | Towards selective phosphodiesterase 2A (PDE2A) inhibitors: a patent review (2010 - present). <i>Expert Opinion on Therapeutic Patents</i> , 2016, 26, 933-946. | 5.0 | 22 |

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|----|---|-----|-----------|
| 19 | Synthesis of 2,1-Borazaroquinolines and 2,1-Borazaroisoquinolines from Vinyl-Aminopyridines and Potassium Organotrifluoroborates by Microwave-Assisted Heating. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 5221-5229. | 2.4 | 17 |
| 20 | Tau Positron Emission Tomography (PET) Imaging: Past, Present, and Future. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 4365-4382. | 6.4 | 88 |
| 21 | Pyrido[4,3- <i>e</i>][1,2,4]triazolo[4,3- <i>a</i>]pyrazines as Selective, Brain Penetrant Phosphodiesterase 2 (PDE2) Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2015, 6, 282-286. | 2.8 | 49 |
| 22 | 1,4-Oxazine $\hat{2}$ -Secretase 1 (BACE1) Inhibitors: From Hit Generation to Orally Bioavailable Brain Penetrant Leads. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 8216-8235. | 6.4 | 67 |
| 23 | Benzazaborinines as Novel Bioisosteric Replacements of Naphthalene: Propranolol as an Example. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 9287-9295. | 6.4 | 62 |
| 24 | Structure-Based Design of a Potent, Selective, and Brain Penetrating PDE2 Inhibitor with Demonstrated Target Engagement. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 1049-1053. | 2.8 | 41 |
| 25 | Regioselective Preparation of 3-Alkoxy-4,5-difluoroanilines by S_N . <i>European Journal of Organic Chemistry</i> , 2012, 2012, 7048-7052. | 2.4 | 7 |
| 26 | Rational design and synthesis of aminopiperazinones as $\hat{2}$ -secretase (BACE) inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 7255-7260. | 2.2 | 44 |
| 27 | Microwave-assisted N-debenzylation of amides with triflic acid. <i>Tetrahedron Letters</i> , 2010, 51, 4815-4818. | 1.4 | 27 |