

Zhude Tu

List of Publications by Citations

Source: <https://exaly.com/author-pdf/7336679/zhude-tu-publications-by-citations.pdf>

Version: 2024-04-29

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

116
papers

3,029
citations

33
h-index

50
g-index

119
ext. papers

3,377
ext. citations

4.7
avg, IF

4.84
L-index

#	Paper	IF	Citations
116	Identification of the PGRMC1 protein complex as the putative sigma-2 receptor binding site. <i>Nature Communications</i> , 2011 , 2, 380	17.4	245
115	Design, synthesis, and evaluation of multitarget-directed resveratrol derivatives for the treatment of Alzheimer's disease. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 5843-59	8.3	180
114	Synthesis and evaluation of multi-target-directed ligands against Alzheimer's disease based on the fusion of donepezil and ebselen. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 9089-99	8.3	125
113	Multitarget-directed benzylideneindanone derivatives: anti- β -amyloid (A β) aggregation, antioxidant, metal chelation, and monoamine oxidase B (MAO-B) inhibition properties against Alzheimer's disease. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 8483-92	8.3	120
112	Fluorine-18-labeled benzamide analogues for imaging the sigma2 receptor status of solid tumors with positron emission tomography. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 3194-204	8.3	95
111	Synthesis and in vitro binding of N-phenyl piperazine analogs as potential dopamine D3 receptor ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2005 , 13, 77-87	3.4	88
110	N-benzylisatin sulfonamide analogues as potent caspase-3 inhibitors: synthesis, in vitro activity, and molecular modeling studies. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 7637-47	8.3	84
109	Binding of the radioligand SIL23 to β -synuclein fibrils in Parkinson disease brain tissue establishes feasibility and screening approaches for developing a Parkinson disease imaging agent. <i>PLoS ONE</i> , 2013 , 8, e55031	3.7	75
108	Benzenediol-berberine hybrids: multifunctional agents for Alzheimer's disease. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 7228-35	3.4	67
107	Design, Synthesis, and Characterization of 3-(Benzylidene)indolin-2-one Derivatives as Ligands for β -Synuclein Fibrils. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 6002-17	8.3	63
106	The novel sigma-2 receptor ligand SW43 stabilizes pancreas cancer progression in combination with gemcitabine. <i>Molecular Cancer</i> , 2010 , 9, 298	42.1	62
105	Carbon-11 labeled sigma2 receptor ligands for imaging breast cancer. <i>Nuclear Medicine and Biology</i> , 2005 , 32, 423-30	2.1	62
104	Inhibition of cholinesterase activity and amyloid aggregation by berberine-phenyl-benzoheterocyclic and tacrine-phenyl-benzoheterocyclic hybrids. <i>Bioorganic and Medicinal Chemistry</i> , 2012 , 20, 3038-48	3.4	58
103	[3H]N-[4-(3,4-dihydro-6,7-dimethoxyisoquinolin-2(1H)-yl)butyl]-2-methoxy-5-methylbenzamide: a novel sigma-2 receptor probe. <i>European Journal of Pharmacology</i> , 2005 , 525, 8-17	5.3	53
102	Radiosynthesis and in vivo evaluation of [11C]MP-10 as a PET probe for imaging PDE10A in rodent and non-human primate brain. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 1666-73	3.4	52
101	Carbon-11 labeled papaverine as a PET tracer for imaging PDE10A: radiosynthesis, in vitro and in vivo evaluation. <i>Nuclear Medicine and Biology</i> , 2010 , 37, 509-16	2.1	47
100	Synthesis and biological evaluation of a new series of ebselen derivatives as glutathione peroxidase (GPx) mimics and cholinesterase inhibitors against Alzheimer's disease. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 1355-61	3.4	45

99	O-Hydroxyl- or o-amino benzylamine-tacrine hybrids: multifunctional biometals chelators, antioxidants, and inhibitors of cholinesterase activity and amyloid- β aggregation. <i>Bioorganic and Medicinal Chemistry</i> , 2012 , 20, 5884-92	3.4	43
98	Synthesis and in vitro and in vivo evaluation of ^{18}F -labeled positron emission tomography (PET) ligands for imaging the vesicular acetylcholine transporter. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 1358-69	8.3	43
97	Synthesis and biological evaluation of a new series of berberine derivatives as dual inhibitors of acetylcholinesterase and butyrylcholinesterase. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 4475-84	3.4	42
96	Pharmacologic characterizations of a P2X7 receptor-specific radioligand, ^{11}C GSK1482160 for neuroinflammatory response. <i>Nuclear Medicine Communications</i> , 2017 , 38, 372-382	1.6	40
95	Characterization of a novel iodinated sigma-2 receptor ligand as a cell proliferation marker. <i>Nuclear Medicine and Biology</i> , 2006 , 33, 203-9	2.1	40
94	Synthesis and in vitro evaluation of β synuclein ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2012 , 20, 4625-34	3.4	38
93	Synthesis and in vivo evaluation of ^{11}C PJ34, a potential radiotracer for imaging the role of PARP-1 in necrosis. <i>Nuclear Medicine and Biology</i> , 2005 , 32, 437-43	2.1	38
92	Quantitative receptor-based imaging of tumor proliferation with the sigma-2 ligand ^{18}F ISO-1. <i>PLoS ONE</i> , 2013 , 8, e74188	3.7	38
91	Novel tacrine-ebesen hybrids with improved cholinesterase inhibitory, hydrogen peroxide and peroxynitrite scavenging activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 6737-42	2.9	37
90	Radiosynthesis and Evaluation of Two PET Radioligands for Imaging β Synuclein. <i>Applied Sciences (Switzerland)</i> , 2014 , 4, 66-78	2.6	37
89	Endogenous dopamine (DA) competes with the binding of a radiolabeled D_1 receptor partial agonist in vivo: a positron emission tomography study. <i>Synapse</i> , 2011 , 65, 724-32	2.4	36
88	Synthesis and in vivo evaluation of 2 high-affinity ^{76}Br -labeled sigma2-receptor ligands. <i>Journal of Nuclear Medicine</i> , 2006 , 47, 1041-8	8.9	36
87	Synthesis and pharmacological evaluation of fluorine-containing D_1 dopamine receptor ligands. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 1555-64	8.3	35
86	Synthesis and characterization of selective dopamine D_2 receptor antagonists. <i>Bioorganic and Medicinal Chemistry</i> , 2006 , 14, 815-25	3.4	34
85	Current status of the development of PET radiotracers for imaging alpha synuclein aggregates in Lewy bodies and Lewy neurites. <i>Clinical and Translational Imaging</i> , 2017 , 5, 3-14	2	33
84	^3H 4-(dimethylamino)-N-(4-(4-(2-methoxyphenyl)piperazin-1-yl) butyl)benzamide: a selective radioligand for dopamine D_3 receptors. II. Quantitative analysis of dopamine D_3 and D_2 receptor density ratio in the caudate-putamen. <i>Synapse</i> , 2010 , 64, 449-59	2.4	33
83	Functional assays to define agonists and antagonists of the sigma-2 receptor. <i>Analytical Biochemistry</i> , 2014 , 448, 68-74	3.1	31
82	Berberine derivatives, with substituted amino groups linked at the 9-position, as inhibitors of acetylcholinesterase/butyrylcholinesterase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 6649-52	2.9	30

81	PET Imaging Study of S1PR1 Expression in a Rat Model of Multiple Sclerosis. <i>Molecular Imaging and Biology</i> , 2016 , 18, 724-32	3.8	28
80	Synthesis and in vitro characterization of a P2X7 radioligand [¹¹ C]TZ6019 and its response to neuroinflammation in a mouse model of Alzheimer disease. <i>European Journal of Pharmacology</i> , 2018 , 820, 8-17	5.3	27
79	Radiosynthesis and biological evaluation of a promising sigma(2)-receptor ligand radiolabeled with fluorine-18 or iodine-125 as a PET/SPECT probe for imaging breast cancer. <i>Applied Radiation and Isotopes</i> , 2010 , 68, 2268-73	1.7	27
78	[³ H]4-(Dimethylamino)-N-[4-(4-(2-methoxyphenyl)piperazin-1-yl)butyl]benzamide, a selective radioligand for dopamine D(3) receptors. I. In vitro characterization. <i>Synapse</i> , 2009 , 63, 717-28	2.4	26
77	The Ru-catalyzed enantioselective preparation of chiral halohydrins and their application in the synthesis of (R)-clorprenaline and (S)-sotalol. <i>Tetrahedron: Asymmetry</i> , 2011 , 22, 722-727		26
76	A promising carbon-11-labeled sphingosine-1-phosphate receptor 1-specific PET tracer for imaging vascular injury. <i>Journal of Nuclear Cardiology</i> , 2017 , 24, 558-570	2.1	25
75	Synthesis and biological characterization of a promising F-18 PET tracer for vesicular acetylcholine transporter. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 4699-4709	3.4	25
74	Synthesis and structure-activity relationship studies of conformationally flexible tetrahydroisoquinolyl triazole carboxamide and triazole substituted benzamide analogues as α receptor ligands. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 4239-51	8.3	24
73	Heteroaromatic and aniline derivatives of piperidines as potent ligands for vesicular acetylcholine transporter. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 6216-33	8.3	23
72	Neuroprotective effects of high affinity β receptor selective compounds. <i>Brain Research</i> , 2012 , 1441, 17-26	3.7	22
71	Synthesis and in vitro biological evaluation of carbonyl group-containing inhibitors of vesicular acetylcholine transporter. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 2825-35	8.3	22
70	Alpha Synuclein Fibrils Contain Multiple Binding Sites for Small Molecules. <i>ACS Chemical Neuroscience</i> , 2018 , 9, 2521-2527	5.7	22
69	Synthesis of Fluorine-Containing Phosphodiesterase 10A (PDE10A) Inhibitors and the In Vivo Evaluation of F-18 Labeled PDE10A PET Tracers in Rodent and Nonhuman Primate. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 8584-600	8.3	21
68	2-(2-Piperidyl)- and 2-(2-pyrrolidyl)chromans as nicotine agonists: synthesis and preliminary pharmacological characterization. <i>Journal of Medicinal Chemistry</i> , 2001 , 44, 4704-15	8.3	19
67	Radiosyntheses and in vivo evaluation of carbon-11 PET tracers for PDE10A in the brain of rodent and nonhuman primate. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 2648-54	3.4	18
66	Hybrids consisting of the pharmacophores of salmeterol and roflumilast or phthalazinone: dual β adrenoceptor agonists-PDE4 inhibitors for the treatment of COPD. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 1548-52	2.9	18
65	Design, Synthesis, and In Vitro and In Vivo Evaluation of an (¹⁸ F)-Labeled Sphingosine 1-Phosphate Receptor 1 (S1P1) PET Tracer. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 6201-20	8.3	17
64	Synthesis and in vitro characterization of cinnoline and benzimidazole analogues as phosphodiesterase 10A inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 919-24	2.9	16

63	Chalcones and Five-Membered Heterocyclic Isosteres Bind to Alpha Synuclein Fibrils in Vitro. <i>ACS Omega</i> , 2018 , 3, 4486-4493	3.9	16
62	Synthesis and evaluation of 15-(4-(2-[¹⁸ F]Fluoroethoxy)phenyl)pentadecanoic acid: a potential PET tracer for studying myocardial fatty acid metabolism. <i>Bioconjugate Chemistry</i> , 2010 , 21, 2313-9	6.3	16
61	Preclinical evaluation of a promising C-11 labeled PET tracer for imaging phosphodiesterase 10A in the brain of living subject. <i>NeuroImage</i> , 2015 , 121, 253-62	7.9	15
60	A potent and selective C-11 labeled PET tracer for imaging sphingosine-1-phosphate receptor 2 in the CNS demonstrates sexually dimorphic expression. <i>Organic and Biomolecular Chemistry</i> , 2015 , 13, 7928-39	3.9	15
59	A practical process for the preparation of [(32)P]S1P and binding assay for S1P receptor ligands. <i>Applied Radiation and Isotopes</i> , 2015 , 102, 5-9	1.7	15
58	An efficient and promising method to prepare Ladostigil (TV3326) via asymmetric transfer hydrogenation catalyzed by RuCl ₃ -DPEN in an HCOONa/H ₂ O/surfactant system. <i>Tetrahedron: Asymmetry</i> , 2012 , 23, 333-338		15
57	Synthesis and biological evaluation of pyrazole group-containing analogues for PDE10A. <i>MedChemComm</i> , 2013 , 4, 443-449	5	15
56	Synthesis and evaluation of in vitro bioactivity for vesicular acetylcholine transporter inhibitors containing two carbonyl groups. <i>Bioorganic and Medicinal Chemistry</i> , 2012 , 20, 4422-9	3.4	15
55	PET Study of Sphingosine-1-Phosphate Receptor 1 Expression in Response to Vascular Inflammation in a Rat Model of Carotid Injury. <i>Molecular Imaging</i> , 2017 , 16, 1536012116689770	3.7	14
54	Glycan array analysis of the antigen repertoire targeted by tumor-binding antibodies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 6839-43	2.9	14
53	Synthesis and in vitro biological evaluation of carbonyl group-containing analogues for α_1 receptors. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 5362-72	8.3	14
52	Syntheses and in vitro evaluation of new S1PR1 compounds and initial evaluation of a lead F-18 radiotracer in rodents. <i>European Journal of Medicinal Chemistry</i> , 2018 , 150, 796-808	6.8	13
51	Effect of cyclosporin A on the uptake of D3-selective PET radiotracers in rat brain. <i>Nuclear Medicine and Biology</i> , 2011 , 38, 725-39	2.1	13
50	Pyrazole Scaffold Synthesis, Functionalization, and Applications in Alzheimer's Disease and Parkinson's Disease Treatment (2011-2020). <i>Molecules</i> , 2021 , 26,	4.8	13
49	Kinetics modeling and occupancy studies of a novel C-11 PET tracer for VACHT in nonhuman primates. <i>Nuclear Medicine and Biology</i> , 2016 , 43, 131-9	2.1	12
48	Computer-assisted designed "selenoxy-chinolin": a new catalytic mechanism of the GPx-like cycle and inhibition of metal-free and metal-associated A β aggregation. <i>Dalton Transactions</i> , 2015 , 44, 20913-25	4.3	11
47	Chemical and enzyme-assisted syntheses of norbuprenorphine-3- β -D-glucuronide. <i>Bioconjugate Chemistry</i> , 2011 , 22, 752-8	6.3	11
46	Upregulated Sphingosine 1-Phosphate Receptor 1 Expression in Human and Murine Atherosclerotic Plaques. <i>Molecular Imaging and Biology</i> , 2018 , 20, 448-456	3.8	11

45	Automated production of [18 F]VAT suitable for clinical PET study of vesicular acetylcholine transporter. <i>Applied Radiation and Isotopes</i> , 2016 , 107, 40-46	1.7	10
44	In vitro and in vivo characterization of two C-11-labeled pet tracers for vesicular acetylcholine transporter. <i>Molecular Imaging and Biology</i> , 2014 , 16, 773-80	3.8	10
43	Preclinical evaluation of the novel monoclonal antibody H6-11 for prostate cancer imaging. <i>Molecular Pharmaceutics</i> , 2013 , 10, 3655-64	5.6	9
42	Radiation dosimetry of [18 F]VAT in nonhuman primates. <i>EJNMMI Research</i> , 2015 , 5, 73	3.6	9
41	Synthesis, radiolabeling and initial in vivo evaluation of [11 C]KSM-01 for imaging PPAR- β receptors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 6233-6	2.9	9
40	Design, synthesis, and in vitro evaluation of quinolinyl analogues for β synuclein aggregation. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018 , 28, 1011-1019	2.9	8
39	Syntheses and radiosyntheses of two carbon-11 labeled potent and selective radioligands for imaging vesicular acetylcholine transporter. <i>Molecular Imaging and Biology</i> , 2014 , 16, 765-72	3.8	8
38	Positron emission tomography imaging of dopamine D2 receptors using a highly selective radiolabeled D2 receptor partial agonist. <i>NeuroImage</i> , 2013 , 71, 168-74	7.9	8
37	Synthesis and in vitro evaluation of new analogues as inhibitors for phosphodiesterase 10A. <i>European Journal of Medicinal Chemistry</i> , 2011 , 46, 3986-95	6.8	8
36	C. Elegans Fatty Acid Two-Hydroxylase Regulates Intestinal Homeostasis by Affecting Heptadecenoic Acid Production. <i>Cellular Physiology and Biochemistry</i> , 2018 , 49, 947-960	3.9	8
35	Kinetic modeling of [18 F]VAT, a novel radioligand for positron emission tomography imaging vesicular acetylcholine transporter in non-human primate brain. <i>Journal of Neurochemistry</i> , 2018 , 144, 791-804	6	7
34	Design and synthesis of pyrazolopyridine derivatives as sphingosine 1-phosphate receptor 2 ligands. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018 , 28, 488-496	2.9	7
33	Development of a carbon-11 PET radiotracer for imaging TRPC5 in the brain. <i>Organic and Biomolecular Chemistry</i> , 2019 , 17, 5586-5594	3.9	6
32	In vitro and ex vivo characterization of (-)-TZ659 as a ligand for imaging the vesicular acetylcholine transporter. <i>European Journal of Pharmacology</i> , 2015 , 752, 18-25	5.3	6
31	In vivo Characterization of Four F-Labeled S1PR1 Tracers for Neuroinflammation. <i>Molecular Imaging and Biology</i> , 2020 , 22, 1362-1369	3.8	6
30	Design, synthesis, and in vitro bioactivity evaluation of fluorine-containing analogues for sphingosine-1-phosphate 2 receptor. <i>Bioorganic and Medicinal Chemistry</i> , 2019 , 27, 3619-3631	3.4	6
29	Absorbed radiation dosimetry of the D-specific PET radioligand [18 F]FluorTriopride estimated using rodent and nonhuman primate. <i>American Journal of Nuclear Medicine and Molecular Imaging</i> , 2016 , 6, 301-309	2.2	6
28	Syntheses and in vitro biological evaluation of S1PR1 ligands and PET studies of four F-18 labeled radiotracers in the brain of nonhuman primates. <i>Organic and Biomolecular Chemistry</i> , 2018 , 16, 9171-9184	3.9	6

27	Synthesis, resolution, and in vitro evaluation of three vesicular acetylcholine transporter ligands and evaluation of the lead fluorine-18 radioligand in a nonhuman primate. <i>Organic and Biomolecular Chemistry</i> , 2017 , 15, 5197-5209	3.9	5
26	Chiral resolution of serial potent and selective ligands and biological evaluation of (-)-[F]TZ3108 in rodent and the nonhuman primate brain. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 1533-1542	3.4	4
25	Comparison of [C]TZ1964B and [F]MNI659 for PET imaging brain PDE10A in nonhuman primates. <i>Pharmacology Research and Perspectives</i> , 2016 , 4, e00253	3.1	4
24	In Vivo Characterization of Two F-Labeled PDE10A PET Radioligands in Nonhuman Primate Brains. <i>ACS Chemical Neuroscience</i> , 2018 , 9, 1066-1073	5.7	3
23	Exploration of Sulfur-Containing Analogues for Imaging Vesicular Acetylcholine Transporter in the Brain. <i>ChemMedChem</i> , 2018 , 13, 1978-1987	3.7	3
22	Automated production of a sphingosine-1 phosphate receptor 1 (S1P1) PET radiopharmaceutical [C]CS1P1 for human use. <i>Applied Radiation and Isotopes</i> , 2019 , 152, 30-36	1.7	3
21	Automated radiochemical synthesis and biodistribution of [11C]-l-acetylmethadol ([11C]LAAM). <i>Applied Radiation and Isotopes</i> , 2014 , 91, 135-40	1.7	3
20	and Investigation of S1PR1 Expression in the Central Nervous System Using [H]CS1P1 and [C]CS1P1. <i>ACS Chemical Neuroscience</i> , 2021 , 12, 3733-3744	5.7	3
19	Radiosynthesis and evaluation of a novel selective PET ligand. <i>MedChemComm</i> , 2014 , 5, 1669-1677	5	2
18	Structure-activity relationship studies and bioactivity evaluation of 1,2,3-triazole containing analogues as a selective sphingosine kinase-2 inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2020 , 206, 112713	6.8	2
17	4-aryl piperidines and 4-(hydroxyphenyl)piperidines as selective sigma-1 receptor ligands: synthesis, preliminary pharmacological evaluation and computational studies. <i>Chemistry Central Journal</i> , 2016 , 10, 53		2
16	Acute Rodent Tolerability, Toxicity, and Radiation Dosimetry Estimates of the S1P1-Specific Radioligand [C]CS1P1. <i>Molecular Imaging and Biology</i> , 2020 , 22, 285-292	3.8	2
15	Radiosynthesis and evaluation of a fluorine-18 labeled radioligand targeting vesicular acetylcholine transporter. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018 , 28, 3425-3430	2.9	2
14	Baseline Microglial Activation Correlates With Brain Amyloidosis and Longitudinal Cognitive Decline in Alzheimer Disease.. <i>Neurology: Neuroimmunology and Neuroinflammation</i> , 2022 , 9,	9.1	2
13	Synthesis and pharmacological evaluation of indolyl carboxylic amide analogues as D dopamine receptor selective ligands. <i>MedChemComm</i> , 2013 , 4, 1283-1289	5	1
12	Automated radiosynthesis of [11C]morphine for clinical investigation. <i>Applied Radiation and Isotopes</i> , 2011 , 69, 431-5	1.7	1
11	Synthesis and evaluation of highly selective quinazoline-2,4-dione ligands for sphingosine-1-phosphate receptor 2.. <i>RSC Medicinal Chemistry</i> , 2022 , 13, 202-207	3.5	1
10	PET Study of Sphingosine-1-phosphate Receptor 1 Expression in Response to Infection.. <i>Molecular Imaging</i> , 2021 , 2021, 9982020	3.7	1

9	In vitro characterization of [H]VAT in cells, animal and human brain tissues for vesicular acetylcholine transporter. <i>European Journal of Pharmacology</i> , 2021 , 911, 174556	5.3	1
8	The impact of dopamine D-like agonist/antagonist on [F]VAT PET measurement of VAcHT in the brain of nonhuman primates. <i>European Journal of Pharmaceutical Sciences</i> , 2020 , 143, 105152	5.1	1
7	Synthesis and characterization of [I]TZ6544, a promising radioligand for investigating sphingosine-1-phosphate receptor 2. <i>Nuclear Medicine and Biology</i> , 2020 , 88-89, 52-61	2.1	1
6	Synthesis and Characterization of a Specific Iodine-125-Labeled TRPC5 Radioligand. <i>ChemMedChem</i> , 2020 , 15, 1854-1860	3.7	1
5	Spatially constrained kinetic modeling with dual reference tissues improves F-flortaucipir PET in studies of Alzheimer disease. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2021 , 48, 3172-3186 ¹	8.8	1
4	Differential Sphingosine-1-Phosphate Receptor-1 Protein Expression in the Dorsolateral Prefrontal Cortex Between Schizophrenia Type 1 and Type 2.. <i>Frontiers in Psychiatry</i> , 2022 , 13, 827981	5	1
3	Cholinergic imbalance in lumbar spinal cord of a rat model of multiple sclerosis. <i>Journal of Neuroimmunology</i> , 2018 , 318, 29-35	3.5	
2	Synthesis and in vitro evaluation of new TRPV4 ligands and biodistribution study of an C-labeled radiotracer in rodents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020 , 30, 127573	2.9	
1	Radiosynthesis and characterization of a carbon-11 PET tracer for receptor-interacting protein kinase 1.. <i>Nuclear Medicine and Biology</i> , 2022 , 110-111, 18-27	2.1	