Victor W Pike

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	PET radiotracers: crossing the blood–brain barrier and surviving metabolism. Trends in Pharmacological Sciences, 2009, 30, 431-440.	4.0	483
2	Chemistry with [¹⁸ F]Fluoride Ion. European Journal of Organic Chemistry, 2008, 2008, 2853-2873.	1.2	413
3	Mixed-Affinity Binding in Humans with 18-kDa Translocator Protein Ligands. Journal of Nuclear Medicine, 2011, 52, 24-32.	2.8	330
4	In vivo radioligand binding to translocator protein correlates with severity of Alzheimer's disease. Brain, 2013, 136, 2228-2238.	3.7	280
5	Tracer Kinetic Modeling of the 5-HT1AReceptor Ligand [carbonyl-11C]WAY-100635 for PET. NeuroImage, 1998, 8, 426-440.	2.1	267
6	Comparison of [11C]-(R)-PK 11195 and [11C]PBR28, two radioligands for translocator protein (18 kDa) in human and monkey: Implications for positron emission tomographic imaging of this inflammation biomarker. NeuroImage, 2010, 49, 2924-2932.	2.1	237
7	A Genetic Polymorphism for Translocator Protein 18 Kda Affects both <i>in Vitro</i> and <i>in Vivo</i> Radioligand Binding in Human Brain to this Putative Biomarker of Neuroinflammation. Journal of Cerebral Blood Flow and Metabolism, 2013, 33, 53-58.	2.4	207
8	Exquisite delineation of 5-HT1A receptors in human brain with PET and [carbonyl-11C]WAY-100635. European Journal of Pharmacology, 1996, 301, R5-R7.	1.7	204
9	Kinetic analysis in healthy humans of a novel positron emission tomography radioligand to image the peripheral benzodiazepine receptor, a potential biomarker for inflammation. NeuroImage, 2008, 40, 43-52.	2.1	193
10	Cerebellum Can Serve As a Pseudo-Reference Region in Alzheimer Disease to Detect Neuroinflammation Measured with PET Radioligand Binding to Translocator Protein. Journal of Nuclear Medicine, 2015, 56, 701-706.	2.8	183
11	Synthesis and Evaluation in Monkey of Two Sensitive ¹¹ C-Labeled Aryloxyanilide Ligands for Imaging Brain Peripheral Benzodiazepine Receptors In Vivo. Journal of Medicinal Chemistry, 2008, 51, 17-30.	2.9	178
12	Reactions of cyclotron-produced [18F]fluoride with diaryliodonium salts—a novel single-step route to no-carrier-added [18]fluoroarenes. Journal of the Chemical Society Chemical Communications, 1995, , 2215-2216.	2.0	168
13	Synthesis of the enantiomers of [N-methyl-11C]PK 11195 and comparison of their behaviours as radioligands for PK binding sites in rats. Nuclear Medicine and Biology, 1994, 21, 573-581.	0.3	164
14	Consensus nomenclature rules for radiopharmaceutical chemistry — Setting the record straight. Nuclear Medicine and Biology, 2017, 55, v-xi.	0.3	162
15	PET imaging with [11C]PBR28 can localize and quantify upregulated peripheral benzodiazepine receptors associated with cerebral ischemia in rat. Neuroscience Letters, 2007, 411, 200-205.	1.0	158
16	The "Specific―P-Glycoprotein Inhibitor Tariquidar Is Also a Substrate and an Inhibitor for Breast Cancer Resistance Protein (BCRP/ABCG2). ACS Chemical Neuroscience, 2011, 2, 82-89.	1.7	153
17	Fast and High-Yield Microreactor Syntheses of <i>ortho</i> -Substituted [¹⁸ F]Fluoroarenes from Reactions of [¹⁸ F]Fluoride Ion with Diaryliodonium Salts. Journal of Organic Chemistry, 2010, 75, 3332-3338.	1.7	149
18	Considerations in the Development of Reversibly Binding PET Radioligands for Brain Imaging. Current Medicinal Chemistry, 2016, 23, 1818-1869.	1.2	149

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19	¹¹ C-ER176, a Radioligand for 18-kDa Translocator Protein, Has Adequate Sensitivity to Robustly Image All Three Affinity Genotypes in Human Brain. Journal of Nuclear Medicine, 2017, 58, 320-325.	2.8	146
20	Persistent Dopamine Functions of Neurons Derived from Embryonic Stem Cells in a Rodent Model of Parkinson Disease. Stem Cells, 2007, 25, 918-928.	1.4	139
21	5â€HT radioligands for human brain imaging with PET and SPECT. Medicinal Research Reviews, 2013, 33, 54-111.	5.0	138
22	Radiation Dosimetry and Biodistribution in Monkey and Man of 11C-PBR28: A PET Radioligand to Image Inflammation. Journal of Nuclear Medicine, 2007, 48, 2072-2079.	2.8	136
23	11C-PBR28 binding to translocator protein increases with progression of Alzheimer's disease. Neurobiology of Aging, 2016, 44, 53-61.	1.5	135
24	PET imaging of the dopamine transporter with 18F-FECNT: a polar radiometabolite confounds brain radioligand measurements. Journal of Nuclear Medicine, 2006, 47, 520-7.	2.8	135
25	P-Glycoprotein Function at the Blood–Brain Barrier in Humans Can Be Quantified with the Substrate Radiotracer ¹¹ C- <i>N-Desmethyl</i> Loperamide. Journal of Nuclear Medicine, 2010, 51, 559-566.	2.8	128
26	Recommendations for fluorine-18 production. International Journal of Radiation Applications and Instrumentation Part A, Applied Radiation and Isotopes, 1991, 42, 749-762.	0.5	126
27	Radioligand Development for PET Imaging of β-Amyloid (Aβ)-Current Status. Current Medicinal Chemistry, 2007, 14, 19-52.	1.2	126
28	Brain and whole-body imaging in nonhuman primates of [11C]PBR28, a promising PET radioligand for peripheral benzodiazepine receptors. NeuroImage, 2008, 39, 1289-1298.	2.1	126
29	Neuroinflammation in Temporal Lobe Epilepsy Measured Using Positron Emission Tomographic Imaging of Translocator Protein. JAMA Neurology, 2015, 72, 882.	4.5	126
30	Remotely-controlled production of the 5-HT1A receptor radioligand, [carbonyl-11C]WAY-100635, via 11C-carboxylation of an immobilized Grignard reagent. Journal of Labelled Compounds and Radiopharmaceuticals, 1996, 38, 941-953.	0.5	117
31	New synthesis of diaryliodonium sulfonates from arylboronic acids. Tetrahedron Letters, 2000, 41, 5393-5396.	0.7	115
32	Imaging the GABA-Benzodiazepine Receptor Subtype Containing the α5-Subunit In Vivo with [11C]Ro15 4513 Positron Emission Tomography. Journal of Cerebral Blood Flow and Metabolism, 2002, 22, 878-889.	2.4	113
33	The Inhibitor Ko143 Is Not Specific for ABCG2. Journal of Pharmacology and Experimental Therapeutics, 2015, 354, 384-393.	1.3	113
34	The synthesis of [18F]fluoroarenes from the reaction of cyclotron-produced [18F]fluoride ion with diaryliodonium salts. Journal of the Chemical Society Perkin Transactions 1, 1998, , 2043-2046.	0.9	108
35	PET evaluation of novel radiofluorinated reboxetine analogs as norepinephrine transporter probes in the monkey brain. Synapse, 2004, 53, 57-67.	0.6	105
36	Small effect of dopamine release and no effect of dopamine depletion on [¹⁸ F]fallypride binding in healthy humans. Synapse, 2008, 62, 399-408.	0.6	104

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37	First delineation of 5-HT1A receptors in human brain with PET and [11C]WAY-100635. European Journal of Pharmacology, 1995, 283, R1-R3.	1.7	103
38	Pre- and post-synaptic dopamine imaging and its relation with frontostriatal cognitive function in Parkinson disease: PET studies with [11C]NNC 112 and [18F]FDOPA. Psychiatry Research - Neuroimaging, 2008, 163, 171-182.	0.9	102
39	Syntheses of 11C- and 18F-labeled carboxylic esters within a hydrodynamically-driven micro-reactor. Lab on A Chip, 2004, 4, 523.	3.1	101
40	Characterization of the radioactive metabolites of the 5-HT1A receptor radioligand, [O-methl-11C]WAY-100635, in monkey and human plasma by HPLC: Comparison of the behaviour of an identified radioactive metabolite with parent radioligand in monkey using PET. Nuclear Medicine and Biology, 1996, 23, 627-634.	0.3	98
41	Translocator Protein PET Imaging for Glial Activation in Multiple Sclerosis. Journal of NeuroImmune Pharmacology, 2011, 6, 354-361.	2.1	98
42	Cu ^I â€Catalyzed ¹¹ Câ€Carboxylation of Boronic Acid Esters: A Rapid and Convenient Entry to ¹¹ Câ€Labeled Carboxylic Acids, Esters, and Amides. Angewandte Chemie - International Edition, 2012, 51, 2698-2702.	7.2	96
43	Characterisation of the Appearance of Radioactive Metabolites in Monkey and Human Plasma from the 5-HT1A Receptor Radioligand, [carbonyl-11C]WAY-100635—Explanation of High Signal Contrast in PET and an Aid to Biomathematical Modelling. Nuclear Medicine and Biology, 1998, 25, 215-223.	0.3	91
44	Synthesis and Evaluation of Two 18F-Labeled 6-Iodo-2-(4â€~-N,N-dimethylamino)phenylimidazo[1,2-a]pyridine Derivatives as Prospective Radioligands for β-Amyloid in Alzheimer's Disease. Journal of Medicinal Chemistry, 2004, 47, 2208-2218.	2.9	91
45	Imaging and Quantitation of Cannabinoid CB ₁ Receptors in Human and Monkey Brains Using ¹⁸ F-Labeled Inverse Agonist Radioligands. Journal of Nuclear Medicine, 2010, 51, 112-120.	2.8	91
46	Synthesis and Evaluation of Translocator 18 kDa Protein (TSPO) Positron Emission Tomography (PET) Radioligands with Low Binding Sensitivity to Human Single Nucleotide Polymorphism rs6971. ACS Chemical Neuroscience, 2014, 5, 963-971.	1.7	91
47	Increased In Vivo Expression of an Inflammatory Marker in Temporal Lobe Epilepsy. Journal of Nuclear Medicine, 2012, 53, 234-240.	2.8	90
48	Regional cerebral opioid receptor studies with [11C]diprenorphine in normal volunteers. Journal of Neuroscience Methods, 1988, 23, 121-129.	1.3	89
49	Fluoridation of heteroaromatic iodonium salts—experimental evidence supporting theoretical prediction of the selectivity of the process. Chemical Communications, 2000, , 649-650.	2.2	89
50	¹¹ C-Loperamide and Its <i>N</i> - <i>Desmethyl</i> Radiometabolite Are Avid Substrates for Brain Permeability-Glycoprotein Efflux. Journal of Nuclear Medicine, 2008, 49, 649-656.	2.8	88
51	Quantitation of cannabinoid CB1 receptors in healthy human brain using positron emission tomography and an inverse agonist radioligand. NeuroImage, 2009, 48, 362-370.	2.1	86
52	Single-Step High-Yield Radiosynthesis and Evaluation of a Sensitive ¹⁸ F-Labeled Ligand for Imaging Brain Peripheral Benzodiazepine Receptors with PET. Journal of Medicinal Chemistry, 2009, 52, 688-699.	2.9	85
53	Positron-emitting radioligands for studies in vivo-probes for human psychopharmacology. Journal of Psychopharmacology, 1993, 7, 139-158.	2.0	81
54	Effect of 5-HT on binding of [11C] WAY 100635 to 5-HT1A receptors in rat brain, assessed using in vivo microdialysis and PET after fenfluramine. Synapse, 2001, 41, 150-159.	0.6	80

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55	Synthesis and Evaluation of [<i>N-</i> methyl- ¹¹ C] <i>N</i> -Desmethyl-loperamide as a New and Improved PET Radiotracer for Imaging P-gp Function. Journal of Medicinal Chemistry, 2008, 51, 6034-6043.	2.9	80
56	The PET Radioligand [11C]MePPEP Binds Reversibly and with High Specific Signal to Cannabinoid CB1 Receptors in Nonhuman Primate Brain. Neuropsychopharmacology, 2008, 33, 259-269.	2.8	80
57	Comparison of four 11C-labeled PET ligands to quantify translocator protein 18ÂkDa (TSPO) in human brain: (R)-PK11195, PBR28, DPA-713, and ER176—based on recent publications that measured specific-to-non-displaceable ratios. EJNMMI Research, 2017, 7, 84.	1.1	80
58	D2 dopamine receptor internalization prolongs the decrease of radioligand binding after amphetamine: A PET study in a receptor internalization-deficient mouse model. NeuroImage, 2010, 50, 1402-1407.	2.1	77
59	Quantification of Translocator Protein (18 kDa) in the Human Brain with PET and a Novel Radioligand, ¹⁸ F-PBR06. Journal of Nuclear Medicine, 2009, 50, 1047-1053.	2.8	75
60	Specific in vivo binding to the norepinephrine transporter demonstrated with the PET radioligand, (S,S)-[11C]MeNER. Nuclear Medicine and Biology, 2003, 30, 707-714.	0.3	74
61	Synthesis of [18F]Fallypride in a Micro-Reactor: Rapid Optimization and Multiple-Production in Small Doses for Micro-PET Studies. Current Radiopharmaceuticals, 2009, 2, 49-55.	0.3	74
62	Synthesis and Simple 18F-Labeling of 3-Fluoro-5-(2-(2-(fluoromethyl)thiazol-4-yl)ethynyl)benzonitrile as a High Affinity Radioligand for Imaging Monkey Brain Metabotropic Glutamate Subtype-5 Receptors with Positron Emission Tomography. Journal of Medicinal Chemistry, 2007, 50, 3256-3266.	2.9	72
63	Facile synthesis of substituted diaryliodonium tosylates by treatment of aryltributylstannanes with Koser's reagent. Journal of the Chemical Society Perkin Transactions 1, 1999, , 245-248.	0.9	69
64	Effect of a P-glycoprotein inhibitor, cyclosporin A, on the disposition in rodent brain and blood of the 5-HT1A receptor radioligand, [11C](R)-(––)-RWAY. Synapse, 2007, 61, 96-105.	0.6	69
65	Synthesis, Ex Vivo Evaluation, and Radiolabeling of Potent 1,5-Diphenylpyrrolidin-2-one Cannabinoid Subtype-1 Receptor Ligands as Candidates for In Vivo Imaging. Journal of Medicinal Chemistry, 2008, 51, 5833-5842.	2.9	69
66	Serotonin-1A receptors in major depression quantified using PET: Controversies, confounds, and recommendations. NeuroImage, 2012, 59, 3243-3251.	2.1	69
67	Kinetic evaluation in nonhuman primates of two new PET ligands for peripheral benzodiazepine receptors in brain. Synapse, 2007, 61, 595-605.	0.6	68
68	Distribution of 5-HT4 receptors in the postmortem human brain—an autoradiographic study using [1251]SB 207710. European Neuropsychopharmacology, 2003, 13, 228-234.	0.3	67
69	A convenient oneâ€pot procedure for the synthesis of 2â€aryl quinazolines using active MnO ₂ as oxidant. Journal of Heterocyclic Chemistry, 2010, 47, 1240-1245.	1.4	67
70	PET-characterization of [carbonyl - 11 C]WAY-100635 binding to 5-HT 1A receptors in the primate brain. Psychopharmacology, 1997, 133, 196-202.	1.5	65
71	Brain and Whole-Body Imaging of Nociceptin/Orphanin FQ Peptide Receptor in Humans Using the PET Ligand ¹¹ C-NOP-1A. Journal of Nuclear Medicine, 2012, 53, 385-392.	2.8	65
72	Post-mortem human brain autoradiography of the norepinephrine transporter using (S,S)-[18F]FMeNER-D2. European Neuropsychopharmacology, 2005, 15, 517-520.	0.3	64

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73	P-Glycoprotein Function at the Blood–Brain Barrier Imaged Using ¹¹ C- <i>N-Desmethyl</i> -Loperamide in Monkeys. Journal of Nuclear Medicine, 2009, 50, 108-115.	2.8	64
74	Synthesis and Evaluation of Radioligands for Imaging Brain Nociceptin/Orphanin FQ Peptide (NOP) Receptors with Positron Emission Tomography. Journal of Medicinal Chemistry, 2011, 54, 2687-2700.	2.9	62
75	Human Brain Imaging and Radiation Dosimetry of ¹¹ C- <i>N</i> - <i>Desmethyl</i> -Loperamide, a PET Radiotracer to Measure the Function of P-Glycoprotein. Journal of Nuclear Medicine, 2009, 50, 807-813.	2.8	61
76	Fluoxetine Administered to Juvenile Monkeys: Effects on the Serotonin Transporter and Behavior. American Journal of Psychiatry, 2014, 171, 323-331.	4.0	61
77	Downregulation of Brain Phosphodiesterase Type IV Measured with 11C-(R)-Rolipram Positron Emission Tomography in Major Depressive Disorder. Biological Psychiatry, 2012, 72, 548-554.	0.7	60
78	Metabotropic Glutamate Subtype 5 Receptors Are Quantified in the Human Brain with a Novel Radioligand for PET. Journal of Nuclear Medicine, 2008, 49, 2042-2048.	2.8	57
79	Influence of alcoholism and cholesterol on TSPO binding in brain: PET [11C]PBR28 studies in humans and rodents. Neuropsychopharmacology, 2018, 43, 1832-1839.	2.8	57
80	Synthesis of a [6- Pyridinyl - 18 F]-labelled fluoro derivative of WAY-100635 as a candidate radioligand for brain 5-HT 1A receptor imaging with PET. Bioorganic and Medicinal Chemistry, 2003, 11, 2769-2782.	1.4	56
81	Identification and regional distribution in rat brain of radiometabolites of the dopamine transporter PET radioligand [11C]PE2I. European Journal of Nuclear Medicine and Molecular Imaging, 2007, 34, 667-678.	3.3	55
82	Rapid and Efficient Radiosyntheses of <i>meta</i> â€6ubstituted [¹⁸ F]Fluoroarenes from [¹⁸ F]Fluoride Ion and Diaryliodonium ÂTosylates within a Microreactor. European Journal of Organic Chemistry, 2011, 2011, 4439-4447.	1.2	55
83	[carbonyl - 11 C]Desmethyl-WAY-100635 (DWAY) is a potent and selective radioligand for central 5-HT 1A receptors in vitro and in vivo. European Journal of Nuclear Medicine and Molecular Imaging, 1998, 25, 338-346.	3.3	54
84	Syntheses of mGluR5 PET radioligands through the radiofluorination of diaryliodonium tosylates. Organic and Biomolecular Chemistry, 2011, 9, 6629.	1.5	54
85	Evaluation of [11C]GB67, a novel radioligand for imaging myocardial α1-adrenoceptors with positron emission tomography. European Journal of Nuclear Medicine and Molecular Imaging, 2000, 27, 7-17.	2.2	53
86	Disulfiram Inhibits Defluorination of 18F-FCWAY, Reduces Bone Radioactivity, and Enhances Visualization of Radioligand Binding to Serotonin 5-HT1A Receptors in Human Brain. Journal of Nuclear Medicine, 2007, 48, 1154-1161.	2.8	52
87	¹¹ C-DPA-713 has much greater specific binding to translocator protein 18 kDa (TSPO) in human brain than ¹¹ C-(<i>R</i>)-PK11195. Journal of Cerebral Blood Flow and Metabolism, 2018, 38, 393-403.	2.4	51
88	Brain and Whole-Body Imaging in Rhesus Monkeys of ¹¹ C-NOP-1A, a Promising PET Radioligand for Nociceptin/Orphanin FQ Peptide Receptors. Journal of Nuclear Medicine, 2011, 52, 1638-1645.	2.8	50
89	Lysosomal trapping of a radiolabeled substrate of P-glycoprotein as a mechanism for signal amplification in PET. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 2593-2598.	3.3	50
90	Population-based input function and image-derived input function for [11C](R)-rolipram PET imaging: Methodology, validation and application to the study of major depressive disorder. NeuroImage, 2012, 63, 1532-1541.	2.1	50

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91	Radioligands for the study of brain 5-HT1A receptors in vivo–development of some new analogues of way. Nuclear Medicine and Biology, 2000, 27, 449-455.	0.3	48
92	PET imaging of brain 5-HT1A receptors in rat in vivo with 18F-FCWAY and improvement by successful inhibition of radioligand defluorination with miconazole. Journal of Nuclear Medicine, 2006, 47, 345-53.	2.8	48
93	The preparation of carbon-11 labelled diprenorphine: a new radioligand for the study of the opiate receptor system in vivo. Journal of the Chemical Society Chemical Communications, 1985, , 1423.	2.0	47
94	A facile and regioselective synthesis of rimonabant through an enamine-directed 1,3-dipolar cycloaddition. Tetrahedron Letters, 2008, 49, 2789-2791.	0.7	47
95	Positron emission tomography imaging using an inverse agonist radioligand to assess cannabinoid CB1 receptors in rodents. Neurolmage, 2008, 41, 690-698.	2.1	47
96	Guidelines for the content and format of PET brain data in publications and archives: A consensus paper. Journal of Cerebral Blood Flow and Metabolism, 2020, 40, 1576-1585.	2.4	47
97	Synthesis of functionalised unsymmetrical diaryliodonium salts. Journal of the Chemical Society Perkin Transactions 1, 1997, , 2463-2466.	0.9	44
98	Radiodefluorination of 3-Fluoro-5-(2-(2-[¹⁸ F](fluoromethyl)-thiazol-4-yl)ethynyl)benzonitrile ([¹⁸ F]SP203), a Radioligand for Imaging Brain Metabotropic Glutamate Subtype-5 Receptors with Positron Emission Tomography, Occurs by Glutathionylation in Rat Brain. Journal of Pharmacology and Experimental Therapeutics, 2008, 327, 727-735.	1.3	44
99	Radiofluorination of diaryliodonium tosylates under aqueous–organic and cryptand-free conditions. Organic and Biomolecular Chemistry, 2013, 11, 5094.	1.5	44
100	Increased Permeability-Glycoprotein Inhibition at the Human Blood–Brain Barrier Can Be Safely Achieved by Performing PET During Peak Plasma Concentrations of Tariquidar. Journal of Nuclear Medicine, 2015, 56, 82-87.	2.8	44
101	An ab initio and MNDO-d SCF-MO computational study of stereoelectronic control in extrusion reactions of R2l–F iodine(III) intermediatesâ€. Journal of the Chemical Society Perkin Transactions II, 1999, , 2707-2714.	0.9	43
102	Evaluation of Two Potent and Selective PET Radioligands to Image COX-1 and COX-2 in Rhesus Monkeys. Journal of Nuclear Medicine, 2018, 59, 1907-1912.	2.8	43
103	Biodistribution and Radiation Dosimetry in Humans of a New PET Ligand, ¹⁸ F-PBR06, to Image Translocator Protein (18 kDa). Journal of Nuclear Medicine, 2010, 51, 145-149.	2.8	42
104	Singleâ€6tep Radiosynthesis of " ¹⁸ Fâ€Labeled Click Synthons―from Azideâ€Functionalized Diaryliodonium Salts. European Journal of Organic Chemistry, 2012, 2012, 4541-4547.	1.2	42
105	Single-step syntheses of no-carrier-added functionalized [18F]fluoroarenes as labeling synthons from diaryliodonium salts. Organic and Biomolecular Chemistry, 2013, 11, 6300.	1.5	42
106	Quantification of brain phosphodiesterase 4 in rat with (R)-[11C]Rolipram-PET. NeuroImage, 2005, 26, 1201-1210.	2.1	41
107	Synthesis and structure–activity relationships (SARs) of 1,5-diarylpyrazole cannabinoid type-1 (CB1) receptor ligands for potential use in molecular imaging. Bioorganic and Medicinal Chemistry, 2006, 14, 3712-3720.	1.4	41
108	Synthesis and Structureâ^'Affinity Relationships of New 4-(6-lodo- <i>H</i> -imidazo[1,2- <i>a</i>]pyridin-2-yl)- <i>N</i> -dimethylbenzeneamine Derivatives as Ligands for Human β-Amyloid Plaques. Journal of Medicinal Chemistry, 2007, 50, 4746-4758.	2.9	41

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109	Regiospecific Syntheses of Functionalized Diaryliodonium Tosylates via [Hydroxy(tosyloxy)iodo]arenes Generated in Situ from (Diacetoxyiodo)arenes. Journal of Organic Chemistry, 2012, 77, 1931-1938.	1.7	41
110	On Quantitative Relationships Between Drug-Like Compound Lipophilicity and Plasma Free Fraction in Monkey and Human. Journal of Pharmaceutical Sciences, 2012, 101, 1028-1039.	1.6	41
111	No-carrier-added [18F]fluoroarenes from the radiofluorination of diaryl sulfoxides. Chemical Communications, 2013, 49, 2151.	2.2	41
112	PET Reveals Inflammation around Calcified Taenia solium Granulomas with Perilesional Edema. PLoS ONE, 2013, 8, e74052.	1.1	41
113	[11C]Carbon monoxide: advances in production and application to PET radiotracer development over the past 15 years. EJNMMI Radiopharmacy and Chemistry, 2019, 4, 25.	1.8	41
114	Improved syntheses of the PET radioligands, [11C]FLB 457, [11C]MDL 100907 and [11C]β-CIT-FE, by the use of [11C]methyl triflate. Journal of Labelled Compounds and Radiopharmaceuticals, 1998, 41, 545-556.	0.5	40
115	Synthesis of [18F]xenon difluoride as a radiolabeling reagent from [18F]fluoride ion in a micro-reactor and at production scale. Journal of Fluorine Chemistry, 2010, 131, 1032-1038.	0.9	40
116	<i>N-desmethyl</i> -Loperamide Is Selective for P-Glycoprotein among Three ATP-Binding Cassette Transporters at the Blood-Brain Barrier. Drug Metabolism and Disposition, 2010, 38, 917-922.	1.7	40
117	Hypervalent aryliodine compounds as precursors for radiofluorination. Journal of Labelled Compounds and Radiopharmaceuticals, 2018, 61, 196-227.	0.5	40
118	Image-Derived Input Function for Human Brain Using High Resolution PET Imaging with [11C](R)-rolipram and [11C]PBR28. PLoS ONE, 2011, 6, e17056.	1.1	40
119	The remotely-controlled preparation of a 11C-labelled radiopharmaceutical—[1â^11C]acetate. The International Journal of Applied Radiation and Isotopes, 1984, 35, 623-627.	0.7	39
120	Amyloid Imaging: From Benchtop to Bedside. Current Topics in Developmental Biology, 2005, 70, 171-213.	1.0	39
121	Quantification of serotonin 5-HT1A receptors in monkey brain with [11C](R)-(â^')-RWAY. Synapse, 2006, 60, 510-520.	0.6	39
122	Distinct patterns of increased translocator protein in posterior cortical atrophy and amnestic Alzheimer's disease. Neurobiology of Aging, 2017, 51, 132-140.	1.5	39
123	4 Radioligands for the Study of Brain 5-HT1A Receptors In Vivo. Progress in Medicinal Chemistry, 2001, 38, 189-247.	4.1	37
124	Kinetic analysis in human brain of [11C](R)-rolipram, a positron emission tomographic radioligand to image phosphodiesterase 4: A retest study and use of an image-derived input function. Neurolmage, 2011, 54, 1903-1909.	2.1	36
125	Decreased Neurokinin-1 (Substance P) Receptor Binding in Patients with Panic Disorder: Positron Emission Tomographic Study with [18F]SPA-RQ. Biological Psychiatry, 2009, 66, 94-97.	0.7	35
126	PET measurement of cyclooxygenase-2 using a novel radioligand: upregulation in primate neuroinflammation and first-in-human study. Journal of Neuroinflammation, 2020, 17, 140.	3.1	35

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127	Radioligands for PET studies of central 5-HT receptors and re-uptake sites — Current status. Nuclear Medicine and Biology, 1995, 22, 1011-1018.	0.3	33
128	Palladium(II)-mediated 11C-carbonylative coupling of diaryliodonium salts with organostannanes—a new, mild and rapid synthesis of aryl [11C]ketones. Journal of the Chemical Society, Perkin Transactions 1, 2000, , 1033-1036.	1.3	33
129	<i>N</i> -(4-Cyanotetrahydro-2 <i>H</i> -pyran-4-yl) and <i>N</i> -(1-Cyanocyclohexyl) Derivatives of 1,5-Diarylpyrazole-3-carboxamides Showing High Affinity for 18 kDa Translocator Protein and/or Cannabinoid Receptors. Journal of Medicinal Chemistry, 2011, 54, 2961-2970.	2.9	33
130	Comparison of 18F- and 11C-labeled aryloxyanilide analogs to measure translocator protein in human brain using positron emission tomography. European Journal of Nuclear Medicine and Molecular Imaging, 2011, 38, 352-357.	3.3	33
131	Biodistribution and radiation dosimetry of a positron emission tomographic ligand, 18F-SP203, to image metabotropic glutamate subtype 5 receptors in humans. European Journal of Nuclear Medicine and Molecular Imaging, 2010, 37, 1943-1949.	3.3	32
132	The PET Radioligand ¹⁸ F-FIMX Images and Quantifies Metabotropic Glutamate Receptor 1 in Proportion to the Regional Density of Its Gene Transcript in Human Brain. Journal of Nuclear Medicine, 2016, 57, 242-247.	2.8	32
133	[¹¹ C]Fluoroform, a Breakthrough for Versatile Labeling of PET Radiotracer Trifluoromethyl Groups in High Molar Activity. Chemistry - A European Journal, 2017, 23, 8156-8160.	1.7	32
134	The PET radioligand [carbonyl-(11)C]desmethyl-WAY-100635 binds to 5-HT(1A) receptors and provides a higher radioactive signal than [carbonyl-(11)C]WAY-100635 in the human brain. Journal of Nuclear Medicine, 2002, 43, 292-303.	2.8	32
135	PET Measurement of the In Vivo Affinity of ¹¹ C-(<i>R</i>)-Rolipram and the Density of Its Target, Phosphodiesterase-4, in the Brains of Conscious and Anesthetized Rats. Journal of Nuclear Medicine, 2009, 50, 749-756.	2.8	31
136	Synthesis and Evaluation in Monkey of [18F]4-Fluoro-N-methyl-N-(4-(6-(methylamino)pyrimidin-4-yl)thiazol-2-yl)benzamide ([18F]FIMX): A Promising Radioligand for PET Imaging of Brain Metabotropic Glutamate Receptor 1 (mGluR1). Journal of Medicinal Chemistry, 2013, 56, 9146-9155.	2.9	31
137	Population-Based Input Function Modeling for [18F]FMPEP-d2, an Inverse Agonist Radioligand for Cannabinoid CB1 Receptors: Validation in Clinical Studies. PLoS ONE, 2013, 8, e60231.	1.1	31
138	Rapid palladium-catalyzed cross-coupling in the synthesis of aryl thioethers under microwave conditions. Tetrahedron Letters, 2006, 47, 4449-4452.	0.7	30
139	Automated radiosynthesis of [18F]SPA-RQ for imaging human brain NK1 receptors with PET. Journal of Labelled Compounds and Radiopharmaceuticals, 2006, 49, 17-31.	0.5	30
140	Propofol Decreases In Vivo Binding of ¹¹ C-PBR28 to Translocator Protein (18 kDa) in the Human Brain. Journal of Nuclear Medicine, 2013, 54, 64-69.	2.8	30
141	Xenon Difluoride Exchanges Fluoride under Mild Conditions: A Simple Preparation of [18F]Xenon Difluoride for PET and Mechanistic Studies. Journal of the American Chemical Society, 2001, 123, 1780-1781.	6.6	29
142	Integration of a microwave reactor with synthia to provide a fully automated radiofluorination module. Journal of Labelled Compounds and Radiopharmaceuticals, 2007, 50, 463-465.	0.5	29
143	Retest imaging of [11C]NOP-1A binding to nociceptin/orphanin FQ peptide (NOP) receptors in the brain of healthy humans. NeuroImage, 2014, 87, 89-95.	2.1	29
144	In vitro and in vivo evaluation of 11C-SD5024, a novel PET radioligand for human brain imaging of cannabinoid CB1 receptors. NeuroImage, 2014, 84, 733-741.	2.1	29

#	Article	IF	CITATIONS
145	3-Substituted 1,5-Diaryl-1 <i>H</i> -1,2,4-triazoles as Prospective PET Radioligands for Imaging Brain COX-1 in Monkey. Part 1: Synthesis and Pharmacology. ACS Chemical Neuroscience, 2018, 9, 2610-2619.	1.7	29
146	PET Radionuclide Production. , 1993, , 1-43.		29
147	A simple technique for the automated production of no-carrier-added [1-11C]acetate. Applied Radiation and Isotopes, 1997, 48, 1117-1120.	0.7	28
148	New halogenated [11C]WAY analogues, [11C]6FPWAY and [11C]6BPWAY—Radiosynthesis and assessment as radioligands for the study of brain 5-HT1A receptors in living monkey. Nuclear Medicine and Biology, 2001, 28, 177-185.	0.3	28
149	Occupancy of dopamine D _{2/3} receptors in rat brain by endogenous dopamine measured with the agonist positron emission tomography radioligand [¹¹ C]MNPA. Synapse, 2008, 62, 756-763.	0.6	28
150	Synthesis and Evaluation of N-Methyl and S-Methyl 11C-Labeled 6-Methylthio-2-(4â€2-N,N-dimethylamino)phenylimidazo[1,2-a]pyridines as Radioligands for Imaging β-Amyloid Plaques in Alzheimer's Disease. Journal of Medicinal Chemistry, 2008, 51, 148-158.	2.9	28
151	[<i>carbonyl</i> â€ ¹¹ C]Benzyl acetate: Automated radiosynthesis via Pdâ€mediated [¹¹ C]carbon monoxide chemistry and PET measurement of brain uptake in monkey. Journal of Labelled Compounds and Radiopharmaceuticals, 2010, 53, 548-551.	0.5	28
152	Selective syntheses of no-carrier-added 2- and 3-[18F]fluorohalopyridines through the radiofluorination of halopyridinyl(4′-methoxyphenyl)iodonium tosylates. Chemical Communications, 2012, 48, 9921.	2.2	28
153	Evaluation of a PET Radioligand to Image <i>O</i> -GlcNAcase in Brain and Periphery of Rhesus Monkey and Knock-Out Mouse. Journal of Nuclear Medicine, 2019, 60, 129-134.	2.8	28
154	First-in-human evaluation of [11C]PS13, a novel PET radioligand, to quantify cyclooxygenase-1 in the brain. European Journal of Nuclear Medicine and Molecular Imaging, 2020, 47, 3143-3151.	3.3	27
155	Preparation of [11C]buprenorphine—A potential radioligand for the study of the opiate receptor system in vivo. International Journal of Radiation Applications and Instrumentation Part A, Applied Radiation and Isotopes, 1987, 38, 65-66.	0.5	26
156	Design of new β1-selective adrenoceptor ligands as potential radioligands for in vivo imaging. Bioorganic and Medicinal Chemistry, 2003, 11, 3513-3527.	1.4	26
157	Radiolabeling of a high potency cannabinoid subtypeâ€1 receptor ligand, <i>N</i> â€(4â€fluoroâ€benzyl)â€4â€(3â€(piperidinâ€1â€yl)â€indoleâ€1â€sulfonyl)benzamide (PipISB), with ca fluorineâ€18. Journal of Labelled Compounds and Radiopharmaceuticals, 2008, 51, 146-152.	bomâ€l1	or26
158	Imaging and In Vivo Quantitation of β-Amyloid: An Exemplary Biomarker for Alzheimer's Disease?. Biological Psychiatry, 2006, 59, 940-947.	0.7	25
159	Quantification of serotonin 5-HT1A receptors in humans with [11C](R)-(â^')-RWAY: Radiometabolite(s) likely confound brain measurements. Synapse, 2007, 61, 469-477.	0.6	25
160	Discovery and Labeling of High-Affinity 3,4-Diarylpyrazolines as Candidate Radioligands for In Vivo Imaging of Cannabinoid Subtype-1 (CB ₁) Receptors. Journal of Medicinal Chemistry, 2008, 51, 5608-5616.	2.9	25
161	Evaluation of Novel <i>N</i> ¹ -Methyl-2-phenylindol-3-ylglyoxylamides as a New Chemotype of 18 kDa Translocator Protein-Selective Ligand Suitable for the Development of Positron Emission Tomography Radioligands. Journal of Medicinal Chemistry, 2011, 54, 366-373.	2.9	25
162	S-[1-(2,3-diaminophenoxy)]-3′-(N-t-butylamino)propan-2′-ol-simplified asymmetric synthesis with CD and chiral HPLC analysis. Tetrahedron: Asymmetry, 1992, 3, 539-554.	1.8	24

#	Article	IF	CITATIONS
163	N-Oxide analogs of WAY-100635: new high affinity 5-HT1A receptor antagonists. Bioorganic and Medicinal Chemistry, 2005, 13, 883-893.	1.4	24
164	¹¹ C-CUMI-101, a PET Radioligand, Behaves as a Serotonin 1A Receptor Antagonist and Also Binds to α ₁ Adrenoceptors in Brain. Journal of Nuclear Medicine, 2014, 55, 141-146.	2.8	24
165	An Investigation of (Diacetoxyiodo)arenes as Precursors for Preparing No-Carrier-Added [¹⁸ F]Fluoroarenes from Cyclotron-Produced [¹⁸ F]Fluoride Ion. Journal of Organic Chemistry, 2016, 81, 297-302.	1.7	24
166	3-Substituted 1,5-Diaryl-1 <i>H</i> -1,2,4-triazoles as Prospective PET Radioligands for Imaging Brain COX-1 in Monkey. Part 2: Selection and Evaluation of [¹¹ C]PS13 for Quantitative Imaging. ACS Chemical Neuroscience, 2018, 9, 2620-2627.	1.7	24
167	Rapid mild syntheses of [11C]benzophenones by Pd(0)-catalysed11C-carbonylative coupling of iodoarenes with phenyltributylstannane in DME-water. Journal of Labelled Compounds and Radiopharmaceuticals, 2000, 43, 825-835.	0.5	23
168	Synthesis and initial evaluation of [11C](R)-RWAY in monkey—a new, simply labeled antagonist radioligand for imaging brain 5-HT1A receptors with PET. European Journal of Nuclear Medicine and Molecular Imaging, 2007, 34, 1670-1682.	3.3	23
169	Open letter to journal editors on: International Consensus Radiochemistry Nomenclature Guidelines. Annals of Nuclear Medicine, 2018, 32, 236-238.	1.2	23
170	Decreased Cannabinoid CB1 Receptors in Male Tobacco Smokers Examined With Positron Emission Tomography. Biological Psychiatry, 2018, 84, 715-721.	0.7	23
171	Synthesis of (+)- and (â^')-cis-2-[(1-adamantylamino)-methyl]-1-phenylcyclopropane derivatives as high affinity probes for σ1 and σ2 binding sites. Il Farmaco, 2002, 57, 45-53.	0.9	22
172	Nucleophile Assisting Leaving Groups: A Strategy for Aliphatic 18F-Fluorination. Journal of Organic Chemistry, 2009, 74, 5290-5296.	1.7	22
173	Candidate PET Radioligand Development for Neurofibrillary Tangles: Two Distinct Radioligand Binding Sites Identified in Postmortem Alzheimer's Disease Brain. ACS Chemical Neuroscience, 2016, 7, 897-911.	1.7	22
174	Pd(0)-Mediated 11C-Carbonylation of Aryl(mesityl)iodonium Salts as a Route to [11C]Arylcarboxylic Acids and Derivatives. Journal of Organic Chemistry, 2017, 82, 11925-11932.	1.7	22
175	Substitution-reduction: an alternative process for the [18F]N-(2-fluoroethylation) of anilines. Journal of Labelled Compounds and Radiopharmaceuticals, 2004, 47, 217-232.	0.5	21
176	Stroke Incidentally Identified Using Improved Positron Emission Tomography for Microglial Activation. Archives of Neurology, 2009, 66, 1288-9.	4.9	21
177	Conformational Structure and Energetics of 2â€Methylphenyl(2′â€methoxyphenyl)iodonium Chloride: Evidence for Solution Clusters. Chemistry - A European Journal, 2010, 16, 10418-10423.	1.7	21
178	Synthesis, Structureâ^'Affinity Relationships, and Radiolabeling of Selective High-Affinity 5-HT ₄ Receptor Ligands as Prospective Imaging Probes for Positron Emission Tomography. Journal of Medicinal Chemistry, 2010, 53, 7035-7047.	2.9	21
179	Development of <i>N</i> -Methyl-(2-arylquinolin-4-yl)oxypropanamides as Leads to PET Radioligands for Translocator Protein (18 kDa). Journal of Medicinal Chemistry, 2014, 57, 6240-6251.	2.9	21
180	Comparison of two PET radioligands, [11C]FPEB and [11C]SP203, for quantification of metabotropic glutamate receptor 5 in human brain. Journal of Cerebral Blood Flow and Metabolism, 2017, 37, 2458-2470.	2.4	21

#	Article	IF	CITATIONS
181	PET ligands [¹⁸ F]LSN3316612 and [¹¹ C]LSN3316612 quantify <i>O</i> -linked-β- <i>N</i> -acetyl-glucosamine hydrolase in the brain. Science Translational Medicine, 2020, 12, .	5.8	21
182	Human biodistribution and radiation dosimetry of the tachykinin NK1 antagonist radioligand [18F]SPA-RQ: comparison of thin-slice, bisected, and 2-dimensional planar image analysis. Journal of Nuclear Medicine, 2007, 48, 100-7.	2.8	21
183	Evaluation of [O-methyl - 11 C]RS-15385-197 as a positron emission tomography radioligand for central α 2 -adrenoceptors. European Journal of Nuclear Medicine and Molecular Imaging, 2000, 27, 475-484.	3.3	20
184	Development of Radioligands for Imaging of Brain Norepinephrine Transporters In Vivo with Positron Emission Tomography. Current Topics in Medicinal Chemistry, 2007, 7, 1806-1816.	1.0	20
185	The decrease of dopamine D2/D3 receptor densities in the putamen and nucleus caudatus goes parallel with maintained levels of CB1 cannabinoid receptors in Parkinson's disease: A preliminary autoradiographic study with the selective dopamine D2/D3 antagonist [3H]raclopride and the novel CB1 inverse agonist [125I]SD7015. Brain Research Bulletin, 2012. 87, 504-510.	1.4	20
186	[¹¹ C]deschloroclozapine is an improved PET radioligand for quantifying a human muscarinic DREADD expressed in monkey brain. Journal of Cerebral Blood Flow and Metabolism, 2021, 41, 2571-2582.	2.4	20
187	Efficient regioselective labelling of the CFC alternative 1,1,1,2-tetrafluoroethane (HFC-134a) with fluorine-18. Journal of Fluorine Chemistry, 1995, 70, 279-287.	0.9	19
188	Short and efficient syntheses of analogues of way-100635: new and potent 5-HT1A receptor antagonists. Bioorganic and Medicinal Chemistry, 2001, 9, 695-702.	1.4	19
189	Radioiodinated SB 207710 as a radioligand in vivo: imaging of brain 5-HT 4 receptors with SPET. European Journal of Nuclear Medicine and Molecular Imaging, 2003, 30, 1520-1528.	3.3	19
190	Synthesis of NCA [carbonyl-11C]amides by direct reaction ofin situ generated [11C]carboxymagnesium halides with amines under microwave-enhanced conditions. Journal of Labelled Compounds and Radiopharmaceuticals, 2003, 46, 1249-1259.	0.5	19
191	The Pyridinyl-6 Position of WAY-100635 as a site for radiofluorination—effect on 5-HT1A receptor radioligand behavior in vivo. Molecular Imaging and Biology, 2004, 6, 17-26.	1.3	19
192	In vivo and in vitro measurement of brain phosphodiesterase 4 in rats after antidepressant administration. Synapse, 2007, 61, 78-86.	0.6	19
193	Positive labeling of ischemic myocardium: A new approach in patients with coronary disease. American Journal of Cardiology, 1981, 47, 481.	0.7	18
194	Radiosyntheses and reactivities of novel [18F]2-fluoroethyl arylsulfonates. Journal of Labelled Compounds and Radiopharmaceuticals, 2005, 48, 735-747.	0.5	18
195	Brain and whole-body imaging in nonhuman primates with [11C]MeS-IMPY, a candidate radioligand for β-amyloid plaques. Nuclear Medicine and Biology, 2007, 34, 681-689.	0.3	18
196	Whole-body biodistribution and radiation dosimetry in monkeys and humans of the phosphodiesterase 4 radioligand [11C](R)-rolipram: comparison of two-dimensional planar, bisected and quadrisected image analyses. Nuclear Medicine and Biology, 2008, 35, 493-500.	0.3	18
197	Effects of ketoconazole on the biodistribution and metabolism of [11C]loperamide and [11C]N-desmethyl-loperamide in wild-type and P-gp knockout mice. Nuclear Medicine and Biology, 2010, 37, 335-345.	0.3	18
198	Radiosynthesis and Evaluation of an ¹⁸ F-Labeled Positron Emission Tomography (PET) Radioligand for Brain Histamine Subtype-3 Receptors Based on a Nonimidazole 2-Aminoethylbenzofuran Chemotype. Journal of Medicinal Chemistry, 2012, 55, 2406-2415.	2.9	18

#	Article	IF	CITATIONS
199	Factors That Limit Positron Emission Tomography Imaging of P-Glycoprotein Density at the Blood–Brain Barrier. Molecular Pharmaceutics, 2013, 10, 2222-2229.	2.3	18
200	Neuroinflammation in frontotemporal lobar degeneration revealed by ¹¹ Câ€PBR28 PET. Annals of Clinical and Translational Neurology, 2019, 6, 1327-1331.	1.7	18
201	EfficientO- andN-(β-fluoroethylation)s with NCA[18F]β-fluoroethyl tosylate under microwave-enhanced conditions. Journal of Labelled Compounds and Radiopharmaceuticals, 2004, 47, 289-297.	0.5	17
202	PET imaging of neurokinin-1 receptors with [18F]SPA-RQ in human subjects: Assessment of reference tissue models and their test–retest reproducibility. Synapse, 2007, 61, 242-251.	0.6	17
203	Effects of cAMPâ€dependent protein kinase activator and inhibitor on in vivo rolipram binding to phosphodiesterase 4 in conscious rats. Synapse, 2010, 64, 172-176.	0.6	17
204	Solution Structures of the Prototypical 18 kDa Translocator Protein Ligand, PK 11195, Elucidated with ¹ H/ ¹³ C NMR Spectroscopy and Quantum Chemistry. ACS Chemical Neuroscience, 2012, 3, 325-335.	1.7	17
205	A Gas Phase Route to [18F]fluoroform with Limited Molar Activity Dilution. Scientific Reports, 2019, 9, 14835.	1.6	17
206	Preparation of carbon-11 labelled prazosin, a potent and selective α1-adrenoreceptor antagonist. Journal of Labelled Compounds and Radiopharmaceuticals, 1988, 25, 177-183.	0.5	16
207	Preparation of no-carrier-added [124I]A14-iodoinsulin as a radiotracer for positron emission tomography. Journal of Labelled Compounds and Radiopharmaceuticals, 2001, 44, 465-480.	0.5	16
208	Efficient and Regioselective Halogenations of 2-Amino-1,3-thiazoles with Copper Salts. Journal of Organic Chemistry, 2009, 74, 2578-2580.	1.7	16
209	Dopamine βâ€hydroxylaseâ€deficient mice have normal densities of D ₂ dopamine receptors in the highâ€affinity state based on in vivo PET imaging and in vitro radioligand binding. Synapse, 2010, 64, 699-703.	0.6	16
210	Synthesis and characterization in monkey of [11C]SP203 as a radioligand for imaging brain metabotropic glutamate 5 receptors. European Journal of Nuclear Medicine and Molecular Imaging, 2012, 39, 1949-1958.	3.3	16
211	Evaluation in vitro and in animals of a new 11C-labeled PET radioligand for metabotropic glutamate receptors 1 in brain. European Journal of Nuclear Medicine and Molecular Imaging, 2013, 40, 245-253.	3.3	16
212	Quinuclidine and DABCO Enhance the Radiofluorination of 5‣ubstituted 2â€Halopyridines. European Journal of Organic Chemistry, 2017, 2017, 6593-6603.	1.2	16
213	2-(4-Methylsulfonylphenyl)pyrimidines as Prospective Radioligands for Imaging Cyclooxygenase-2 with PET—Synthesis, Triage, and Radiolabeling. Molecules, 2018, 23, 2850.	1.7	16
214	Building a database for brain 18 kDa translocator protein imaged using [¹¹ C]PBR28 in healthy subjects. Journal of Cerebral Blood Flow and Metabolism, 2019, 39, 1138-1147.	2.4	16
215	Whole-body biodistribution and estimation of radiation-absorbed doses of the dopamine D1 receptor radioligand 11C-NNC 112 in humans. Journal of Nuclear Medicine, 2006, 47, 100-4.	2.8	16
216	An ab initio and MNDO-d SCF–MO computational study of the extrusion reactions of R2I–F iodine(III) via dimeric, trimeric and tetrameric transition states â€. Perkin Transactions II RSC, 2000, , 2158-2161.	1.1	15

#	Article	IF	CITATIONS
217	Development of Novel Amyloid Imaging Agents Based Upon Thioflavin S. Current Alzheimer Research, 2005, 2, 109-114.	0.7	15
218	Investigation of the Metabolites of (S,S)-[11C]MeNER in Humans, Monkeys and Rats. Molecular Imaging and Biology, 2009, 11, 23-30.	1.3	15
219	Evaluation of [¹¹ C]PipISB and [¹⁸ F]PipISB in monkey as candidate radioligands for imaging brain cannabinoid type†receptors in vivo. Synapse, 2009, 63, 22-30.	0.6	15
220	Evaluation of ¹¹ C-NR2B-SMe and Its Enantiomers as PET Radioligands for Imaging the NR2B Subunit Within the NMDA Receptor Complex in Rats. Journal of Nuclear Medicine, 2020, 61, 1212-1220.	2.8	15
221	[¹¹ C]Carbonyl Difluoride—a New and Highly Efficient [¹¹ C]Carbonyl Group Transfer Agent. Angewandte Chemie - International Edition, 2020, 59, 7256-7260.	7.2	15
222	Human Biodistribution and Dosimetry of 11C-CUMI-101, an Agonist Radioligand for Serotonin-1A Receptors in Brain. PLoS ONE, 2011, 6, e25309.	1.1	15
223	Synthesis and Screening in Mice of Fluorine-Containing PET Radioligands for TSPO: Discovery of a Promising ¹⁸ F-Labeled Ligand. Journal of Medicinal Chemistry, 2021, 64, 16731-16745.	2.9	15
224	Semi-automated preparation of a 11C-labelled antibiotic-[N-methyl-11C]Erthromycin A lactobionate. The International Journal of Applied Radiation and Isotopes, 1984, 35, 103-109.	0.7	14
225	Efficient and selective labelling of the CFC alternative, 1,1,1,2-tetrafluoroethane, with 18F in the 1-position. Journal of the Chemical Society Chemical Communications, 1993, , 1064.	2.0	14
226	Propionyl-l-carnitine: Labelling in the N-methyl position with carbon-11 and pharmacokinetic studies in rats. Nuclear Medicine and Biology, 1995, 22, 699-709.	0.3	14
227	Automated chemoenzymatic synthesis of no-carrier-added [carbonyl-11C]propionyl l-carnitine for pharmacokinetic studies. Applied Radiation and Isotopes, 1997, 48, 917-924.	0.7	14
228	Synthesis and first in vivo evaluation of new selective high affinity β1-adrenoceptor radioligands for SPECT based on ICI 89,406. Bioorganic and Medicinal Chemistry, 2004, 12, 4117-4132.	1.4	14
229	Synthesis and PET evaluation of (R)-[S-methyl-11C]thionisoxetine, a candidate radioligand for imaging brain norepinephrine transporters. Journal of Labelled Compounds and Radiopharmaceuticals, 2006, 49, 1007-1019.	0.5	14
230	Synthesis of 11C-labelled (R)-OHDMI and CFMME and their evaluation as candidate radioligands for imaging central norepinephrine transporters with PET. Bioorganic and Medicinal Chemistry, 2007, 15, 616-625.	1.4	14
231	The [18F]2-fluoro-1,3-thiazolyl moiety—an easily-accessible structural motif for prospective molecular imaging radiotracers. Tetrahedron Letters, 2010, 51, 6034-6036.	0.7	14
232	Image-derived input function in PET brain studies. Nuclear Medicine Communications, 2012, 33, 982-989.	0.5	14
233	[11C]Rhodamine-123: Synthesis and biodistribution in rodents. Nuclear Medicine and Biology, 2012, 39, 1128-1136.	0.3	14
234	Synthesis and evaluation of candidate PET radioligands for corticotropin-releasing factor type-1 receptors. Nuclear Medicine and Biology, 2014, 41, 524-535.	0.3	14

#	Article	IF	CITATIONS
235	<i>N</i> â€2-3-(Trifluoromethyl)phenyl Derivatives of <i>N</i> Aryl- <i>N</i> â€2-methylguanidines as Prospective PET Radioligands for the Open Channel of the <i>N</i> -Methyl- <scp>d</scp> -aspartate (NMDA) Receptor: Synthesis and Structure–Affinity Relationships. Journal of Medicinal Chemistry, 2015, 58, 9722-9730.	2.9	14
236	[carbonyl - 11 C]4-Fluoro- N -methyl- N -(4-(6-(methylamino)pyrimidin-4-yl)thiazol-2-yl)benzamide ([11) Tj ETQqC monkey brain. Nuclear Medicine and Biology, 2015, 42, 967-974.	0 0 rgBT 0.3	/Overlock 10 14
237	Exploration of the labeling of [¹¹ C]tubastatin A at the hydroxamic acid site with [¹¹ C]carbon monoxide. Journal of Labelled Compounds and Radiopharmaceuticals, 2016, 59, 9-13.	0.5	14
238	Papain, chymotrypsin and related proteins—a comparative study of their beer chill-proofing abilities and characteristics. Enzyme and Microbial Technology, 1981, 3, 59-63.	1.6	13
239	Kinetic brain analysis and wholeâ€body imaging in monkey of [¹¹ C]MNPA: A dopamine agonist radioligand. Synapse, 2008, 62, 700-709.	0.6	13
240	Quantification of metabotropic glutamate subtype 5 receptors in the brain by an equilibrium method using 18F-SP203. Neurolmage, 2012, 59, 2124-2130.	2.1	13
241	Image-Derived Input Function Derived from a Supervised Clustering Algorithm: Methodology and Validation in a Clinical Protocol Using [11C](R)-Rolipram. PLoS ONE, 2014, 9, e89101.	1.1	13
242	Preparation of a carbon-11 labelled antibiotic, erythromycin A lactobionate. Journal of the Chemical Society Chemical Communications, 1982, , 173.	2.0	12
243	Visualization and characterization of 5-HT receptors and transporters in vivo and in man. Seminars in Neuroscience, 1995, 7, 421-431.	2.3	12
244	Development of central 5-HT2A receptor radioligands for PET: Comparison of [3H]RP 62203 and [3H]SR 46349B kinetics in rat brain. Nuclear Medicine and Biology, 1996, 23, 245-250.	0.3	12
245	Automated radiosynthesis of no-carrier-added [S-fluoromethyl- 18F]fluticasone propionate as a radiotracer for lung deposition studies with PET. Journal of Labelled Compounds and Radiopharmaceuticals, 1997, 39, 567-584.	0.5	12
246	Lipophilic Analogs of Thioflavin S as Novel Amyloid-Imaging Agents. Current Alzheimer Research, 2006, 3, 259-266.	0.7	12
247	Synthesis and in vitro autoradiographic evaluation of a novel high-affinity radioiodinated ligand for imaging brain cannabinoid subtype-1 receptors. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 6209-6212.	1.0	12
248	Preclinical evaluation of an 18F-labelled $\hat{1}^2$ 1-adrenoceptor selective radioligand based on ICI 89,406. Nuclear Medicine and Biology, 2010, 37, 517-526.	0.3	12
249	Rapid Roomâ€Temperature ¹¹ Câ€Methylation of Arylamines with [¹¹ C]Methyl Iodide Promoted by Solid Inorganicâ€Bases in DMF. European Journal of Organic Chemistry, 2012, 2012, 1303-1310.	1.2	12
250	A PET study comparing receptor occupancy by five selective cannabinoid 1 receptor antagonists in non-human primates. Neuropharmacology, 2016, 101, 519-530.	2.0	12
251	Discovery, Radiolabeling, and Evaluation of Subtype-Selective Inhibitors for Positron Emission Tomography Imaging of Brain Phosphodiesterase-4D. ACS Chemical Neuroscience, 2020, 11, 1311-1323.	1.7	12
252	Synthesis of [¹⁸ F]PS13 and Evaluation as a PET Radioligand for Cyclooxygenase-1 in Monkey. ACS Chemical Neuroscience, 2021, 12, 517-530.	1.7	12

#	Article	IF	CITATIONS
253	Syntheses of 2-Amino and 2-Halothiazole Derivatives as High-Affinity Metabotropic Glutamate Receptor Subtype 5 Ligands and Potential Radioligands for in Vivo Imaging. Journal of Medicinal Chemistry, 2011, 54, 901-908.	2.9	11
254	Enhanced nucleophilic fluorination and radiofluorination of organosilanes appended with potassium-chelating leaving groups. Journal of Fluorine Chemistry, 2014, 158, 48-52.	0.9	11
255	Crystal Structures of Diaryliodonium Fluorides and Their Implications for Fluorination Mechanisms. Chemistry - A European Journal, 2017, 23, 4353-4363.	1.7	11
256	Rapid and Efficient Synthesis of [¹¹ C]Trifluoromethylarenes from Primary Aromatic Amines and [¹¹ C]CuCF ₃ . ACS Omega, 2020, 5, 19557-19564.	1.6	11
257	11C-LY2428703, a positron emission tomographic radioligand for the metabotropic glutamate receptor 1, is unsuitable for imaging in monkey and human brains. EJNMMI Research, 2013, 3, 47.	1.1	10
258	18F-FCWAY, a serotonin 1A receptor radioligand, is a substrate for efflux transport at the human blood-brain barrier. NeuroImage, 2016, 138, 134-140.	2.1	10
259	Alternative methods for labeling the 5-HT1A receptor agonist, 1-[2-(4-fluorobenzoylamino)ethyl]-4-(7-methoxynaphthyl)piperazine (S14506), with carbon-11 or fluorine-18. Journal of Labelled Compounds and Radiopharmaceuticals, 2005, 48, 971-981.	0.5	9
260	In vivo binding of protoporphyrin IX to rat translocator protein imaged with positron emission tomography. Synapse, 2010, 64, 649-653.	0.6	9
261	Evaluation of [11C]S14506 and [18F]S14506 in Rat and Monkey as Agonist PET Radioligands for Brain 5-HT1A Receptors. Current Radiopharmaceuticals, 2010, 3, 9-18.	0.3	9
262	Essential Principles and Recent Progress in the Development of TSPO PET Ligands for Neuroinflammation Imaging. Current Medicinal Chemistry, 2022, 29, 4862-4890.	1.2	9
263	A study of the beer chill-proofing behaviour of a water-insoluble papain conjugate of hydrous titanium(IV) oxide-use of hydrous titanium(IV) oxide as a novel chill-proofing agent. Enzyme and Microbial Technology, 1980, 2, 126-132.	1.6	8
264	Novel use of an isotope separator to determine the position of fluorine-18 in labelled 1,1,1,2-tetrafluoroethanes. Organic Mass Spectrometry, 1994, 29, 499-504.	1.3	8
265	Synthesis of 2- and 6-fluoro analogues of threo-3-(3,4-dihydroxyphenyl)serine (2- and) Tj ETQq1 1 0.784314 rg	BT /Overloc	k 10 Tf 50 20
266	Whole-body biodistribution and radiation dosimetry estimates for the PET dopamine transporter probe 18F-FECNT in non-human primates. Nuclear Medicine Communications, 2004, 25, 737-742.	0.5	8
267	Identification of the Transition States in the Inversion of 1,4-Benzodiazepines with the <i>Ab Initio</i> Replica Path Method. Journal of Physical Chemistry A, 2008, 112, 1604-1611.	1.1	8
268	Evaluation in monkey of two candidate PET radioligands, [¹¹ C]RXâ€1 and [¹⁸ F]RXâ€2, for imaging brain 5â€HT ₄ receptors. Synapse, 2014, 68, 613-623.	0.6	8
269	[11 C]AZ10419096 – a full antagonist PET radioligand for imaging brain 5-HT 1B receptors. Nuclear Medicine and Biology, 2017, 54, 34-40.	0.3	8
270	Rapid Syntheses of [¹¹ C]Arylvinyltrifluoromethanes through Treatment of (<i>E</i>)-Arylvinyl(phenyl)iodonium Tosylates with [¹¹ C]Trifluoromethylcopper(I). Organic Letters, 2020, 22, 4574-4578.	2.4	8

#	Article	IF	CITATIONS
271	The Synthesis of Co-polymers with Pendant Functional Groups Arranged in a Predetermined Geometry as Enzyme Models. Biochemical Society Transactions, 1978, 6, 269-271.	1.6	7
272	Synthesis of no-carrier-added [11C]methanesulfonyl chloride as a new labeling agent for PET radiopharmaceutical development. Journal of Labelled Compounds and Radiopharmaceuticals, 2003, 46, 1127-1140.	0.5	7
273	Crown Ether Nucleophilic Catalysts (CENCs): Agents for Enhanced Silicon Radiofluorination. Journal of Organic Chemistry, 2017, 82, 2329-2335.	1.7	7
274	11C-Labeling of Aryl Ketones as Candidate Histamine Subtype-3 Receptor PET Radioligands through Pd(0)-Mediated 11C-Carbonylative Coupling. Molecules, 2017, 22, 792.	1.7	7
275	Synthesis and evaluation of two new candidate high-affinity full agonist PET radioligands for imaging 5-HT1B receptors. Nuclear Medicine and Biology, 2019, 70, 1-13.	0.3	7
276	Status of the â€~consensus nomenclature rules in radiopharmaceutical sciences' initiative. Nuclear Medicine and Biology, 2019, 71, 19-22.	0.3	7
277	Development of a 18F-labeled PET radioligand for imaging 5-HT1B receptors: [18F]AZ10419096. Nuclear Medicine and Biology, 2019, 78-79, 11-16.	0.3	7
278	Syntheses of [¹¹ C]2- and [¹¹ C]3-trifluoromethyl-4-aminopyridine: potential PET radioligands for demyelinating diseases. RSC Medicinal Chemistry, 2020, 11, 1161-1167.	1.7	7
279	The chemistry of labeling heterocycles with carbon-11 or fluorine-18 for biomedical imaging. Advances in Heterocyclic Chemistry, 2020, 132, 241-384.	0.9	7
280	Translation of 11C-labeled tracer synthesis to a CGMP environment as exemplified by [11C]ER176 for PET imaging of human TSPO. Nature Protocols, 2021, 16, 4419-4445.	5.5	7
281	The radiosynthesis of nca [O-methyl-11C]viqualine, through an N-trityl-protected intermediate, as a potential pet radioligand for 5HT re-uptake sites. Journal of Labelled Compounds and Radiopharmaceuticals, 1990, 28, 1341-1350.	0.5	6
282	Avoiding Barriers to PET Radioligand Development: Cellular Assays of Brain Efflux Transporters. Journal of Nuclear Medicine, 2011, 52, 338-340.	2.8	6
283	PET Imaging of Phosphodiesterase-4 Identifies Affected Dysplastic Bone in McCune–Albright Syndrome, a Genetic Mosaic Disorder. Journal of Nuclear Medicine, 2020, 61, 1672-1677.	2.8	6
284	Repurposing ¹¹ C-PS13 for PET Imaging of Cyclooxygenase-1 in Ovarian Cancer Xenograft Mouse Models. Journal of Nuclear Medicine, 2021, 62, 665-668.	2.8	6
285	Cyclooxygenases as Potential PET Imaging Biomarkers to Explore Neuroinflammation in Dementia. Journal of Nuclear Medicine, 2022, 63, 53S-59S.	2.8	6
286	Significant abilities of titanium(IV)-activated glass fibre paper and its papain conjugates to chill-proof beer. Enzyme and Microbial Technology, 1980, 2, 288-294.	1.6	5
287	New N-aryl-N′-(3-(substituted)phenyl)-N′-methylguanidines as leads to potential PET radioligands for imaging the open NMDA receptor. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 225-228. 	1.0	5
288	[<i>Carboxyl</i> â€ ¹¹ C]Labelling of Four Highâ€Affinity cPLA2α Inhibitors and Their Evaluation as Radioligands in Mice by Positron Emission Tomography. ChemMedChem, 2018, 13, 138-146.	1.6	5

#	Article	IF	CITATIONS
289	Open letter to journal editors on: International consensus radiochemistry nomenclature guidelines. Journal of Labelled Compounds and Radiopharmaceuticals, 2018, 61, 402-404.	0.5	5
290	Broad Scope and Highâ€Yield Access to Unsymmetrical Acyclic [¹¹ C]Ureas for Biomedical Imaging from [¹¹ C]Carbonyl Difluoride. Chemistry - A European Journal, 2021, 27, 10369-10376.	1.7	5
291	Low Retention of [S-methyl-11C]MeS-IMPY to β-amyloid Plaques in Patients with Alzheimers Disease. Current Radiopharmaceuticals, 2009, 2, 129-136.	0.3	5
292	The preparation of a carbon-11 labelled analgesic - [N-methyl-11C]meptazinol. Journal of Labelled Compounds and Radiopharmaceuticals, 1987, 24, 1051-1059.	0.5	4
293	Development of new brain imaging agents based upon nocaine–modafinil hybrid monoamine transporter inhibitors. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 3101-3104.	1.0	4
294	Characterization of fast-decaying PET radiotracers solely through LC-MS/MS of constituent radioactive and carrier isotopologues. EJNMMI Research, 2013, 3, 3.	1.1	4
295	T80. Novel PET Radioligands Show That COX-2, but not COX-1, is Induced by Neuroinflammation in Rhesus Macaque. Biological Psychiatry, 2018, 83, S160.	0.7	4
296	[11 C]Carbonyl Difluoride—a New and Highly Efficient [11 C]Carbonyl Group Transfer Agent. Angewandte Chemie, 2020, 132, 7323-7327.	1.6	4
297	Potential for imaging the high-affinity state of the 5-HT1B receptor: a comparison of three PET radioligands with differing intrinsic activity. EJNMMI Research, 2019, 9, 100.	1.1	4
298	Labelling of the CFC-alternative, 2H-heptafluoropropane (HFC 227ea), with fluorine-18. Journal of Fluorine Chemistry, 1995, 75, 67-73.	0.9	3
299	[11C](R)-Rolipram positron emission tomography detects DISC1 inhibition of phosphodiesterase type 4 in live Disc1 locus-impaired mice. Journal of Cerebral Blood Flow and Metabolism, 2019, 39, 1306-1313.	2.4	3
300	Development of a non-radiometric method for measuring the arterial input function of a 11C-labeled PET radiotracer. Scientific Reports, 2020, 10, 17350.	1.6	3
301	In vivo SPECT and ex vivo autoradiographic brain imaging of the novel selective CB1 receptor antagonist radioligand [125I]SD7015 in CB1 knock-out and wildtype mouse. Brain Research Bulletin, 2013, 91, 46-51.	1.4	2
302	Translocator protein ligands based on N -methyl-(quinolin-4-yl)oxypropanamides with properties suitable for PET radioligand development. European Journal of Medicinal Chemistry, 2016, 124, 677-688.	2.6	2
303	[O-methyl-11C]N-(4-(4-(3-Chloro-2-methoxyphenyl)-piperazin-1-yl)butyl)-1H-indole-2-carboxamide ([11C]BAK4-51) Is an Efflux Transporter Substrate and Ineffective for PET Imaging of Brain D3 Receptors in Rodents and Monkey. Molecules, 2018, 23, 2737.	1.7	2
304	Region- and voxel-based quantification in human brain of [18F]LSN3316612, a radioligand for O-GlcNAcase. EJNMMI Research, 2021, 11, 35.	1.1	2
305	Fluorine-18 chemistry in micro-reactors. Journal of Labelled Compounds and Radiopharmaceuticals, 2010, 53, 234-238.	0.5	2
306	Acetylation of <i>N</i> -Heteroaryl Bromides via PdCl ₂ /(<i>o</i> -tolyl) ₃ P Catalyzed Heck Reactions. Synthesis, 2008, 2008, 887-890.	1.2	1

#	Article	IF	CITATIONS
307	On the quantitative relationship between radiotracer lipophilicity and plasma free fraction. NeuroImage, 2010, 52, S221.	2.1	1
308	Front Cover: Quinuclidine and DABCO Enhance the Radiofluorination of 5-Substituted 2-Halopyridines (Eur. J. Org. Chem. 45/2017). European Journal of Organic Chemistry, 2017, 2017, 6575-6575.	1.2	1
309	International Consensus Radiochemistry Nomenclature Guidelines. Radiochimica Acta, 2018, 106, 623-625.	0.5	1
310	Radiosynthesis and Evaluation in Monkey of Three 11C-Labeled 1,5-Diarylpyrazoles as High Potency Candidate PET Radioligands for Cannabinoid Subtype-1 Receptors in Brain. Current Radiopharmaceuticals, 2008, 1, 93-102.	0.3	1
311	Increased Regional Myocardial Glucose Utilisation in Patients with Chronic Stable Angina as Assessed by Positron Emission Tomography (PET). Clinical Science, 1989, 76, 55P-55P.	0.0	0
312	Repurposing [11C]MC1 for PET Imaging of Cyclooxygenase-2 in Colorectal Cancer Xenograft Mouse Models. Molecular Imaging and Biology, 2021, , 1.	1.3	0
313	Tandem Mass Spectrometry as an Independent Method for Corroborating Fluorine-18 Radioactivity Measurements in Positron Emission Tomography. ACS Measurement Science Au, 2022, 2, 370-376.	1.9	0