

# Xavier Langlois

## List of Publications by Year in descending order

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53  
papers

2,685  
citations

186265

28  
h-index

182427

51  
g-index

54  
all docs

54  
docs citations

54  
times ranked

3502  
citing authors

#	ARTICLE	IF	CITATIONS
1	Striatal phosphodiesterase 10A availability is altered secondary to chronic changes in dopamine neurotransmission. <i>EJNMMI Radiopharmacy and Chemistry</i> , 2017, 1, 3.	3.9	13
2	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
3	Preclinical Evaluation of <sup>18</sup> F-JNJ64349311, a Novel PET Tracer for Tau Imaging. <i>Journal of Nuclear Medicine</i> , 2017, 58, 975-981.	5.0	72
4	Evaluation of Small-Animal PET Outcome Measures to Detect Disease Modification Induced by BACE Inhibition in a Transgenic Mouse Model of Alzheimer Disease. <i>Journal of Nuclear Medicine</i> , 2017, 58, 1977-1983.	5.0	24
5	What We Observe In Vivo Is Not Always What We See In Vitro: Development and Validation of <sup>11</sup> C-JNJ-42491293, A Novel Radioligand for mGluR2. <i>Journal of Nuclear Medicine</i> , 2017, 58, 110-116.	5.0	31
6	The Effects of Physiological and Methodological Determinants on <sup>18</sup> F-FDG Mouse Brain Imaging Exemplified in a Double Transgenic Alzheimer Model. <i>Molecular Imaging</i> , 2016, 15, 153601211562491.	1.4	21
7	Metabotropic glutamate receptor 2/3 density and its relation to the hippocampal neuropathology in a model of temporal lobe epilepsy in rats. <i>Epilepsy Research</i> , 2016, 127, 55-59.	1.6	6
8	BiDiFuse: a Fiji plugin for fusing bi-directionally recorded microscopic image volumes. <i>Bioinformatics</i> , 2016, 32, 3691-3693.	4.1	2
9	Longitudinal Characterization of [ <sup>18</sup> F]-FDG and [ <sup>18</sup> F]-AV45 Uptake in the Double Transgenic TASTPM Mouse Model. <i>Journal of Alzheimer's Disease</i> , 2016, 55, 1537-1548.	2.6	15
10	Image Informatics Strategies for Deciphering Neuronal Network Connectivity. <i>Advances in Anatomy, Embryology and Cell Biology</i> , 2016, 219, 123-148.	1.6	5
11	Preclinical evaluation of the antipsychotic potential of the mGlu2-positive allosteric modulator JNJ40411813. <i>Pharmacology Research and Perspectives</i> , 2015, 3, e00097.	2.4	39
12	In vivo molecular neuroimaging of glucose utilization and its association with fibrillar amyloid- $\beta$ load in aged APPS1-21 mice. <i>Alzheimer's Research and Therapy</i> , 2015, 7, 76.	6.2	27
13	Peripheral Administration of Tumor Necrosis Factor-Alpha Induces Neuroinflammation and Sickness but Not Depressive-Like Behavior in Mice. <i>BioMed Research International</i> , 2015, 2015, 1-14.	1.9	50
14	Discovery of VU0409551/JNJ-46778212: An mGlu <sub>5</sub> Positive Allosteric Modulator Clinical Candidate Targeting Schizophrenia. <i>ACS Medicinal Chemistry Letters</i> , 2015, 6, 716-720.	2.8	41
15	Identification of a Novel Orally Bioavailable Phosphodiesterase 10A (PDE10A) Inhibitor with Efficacy in Animal Models of Schizophrenia.. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 978-993.	6.4	16
16	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
17	Pharmacological and pharmacokinetic properties of JNJ40411813, a positive allosteric modulator of the mGlu2 receptor. <i>Pharmacology Research and Perspectives</i> , 2015, 3, e00096.	2.4	32
18	Preclinical Comparison of the Amyloid- $\beta$ Radioligands [ <sup>11</sup> C]Pittsburgh compound B and [ <sup>18</sup> F]florbetaben in Aged APPS1-21 and BRI1-42 Mouse Models of Cerebral Amyloidosis. <i>Molecular Imaging and Biology</i> , 2015, 17, 688-696.	2.6	8

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19	Effect of stress and peripheral immune activation on astrocyte activation in transgenic bioluminescent <i>GLT-1</i> mice. <i>Glia</i> , 2015, 63, 1126-1137.	4.9	22
20	Quantitative $^{18}\text{F}$ PET Imaging of Cerebral Glucose Metabolism and Amyloidosis in the TASTPM Double Transgenic Mouse Model of Alzheimer's Disease. <i>Current Alzheimer Research</i> , 2015, 12, 694-703.	1.4	14
21	Examining dopamine D3 receptor occupancy by antipsychotic drugs via [ $^3\text{H}$ ]7-OH-DPAT <i>ex vivo</i> autoradiography and its cross-validation via c-fos immunohistochemistry in the rat brain. <i>European Journal of Pharmacology</i> , 2014, 740, 669-675.	3.5	9
22	The [ $^{18}\text{F}$ ]FDG $^{18}\text{F}$ PET Readout of a Brain Activation Model to Evaluate the Metabotropic Glutamate Receptor 2 Positive Allosteric Modulator JNJ-42153605. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2014, 350, 375-386.	2.5	12
23	Discovery of a Potent, Selective, and Orally Active Phosphodiesterase 10A Inhibitor for the Potential Treatment of Schizophrenia. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 4196-4212.	6.4	36
24	Pharmacology of JNJ-42314415, a Centrally Active Phosphodiesterase 10A (PDE10A) Inhibitor: A Comparison of PDE10A Inhibitors with D <sub>2</sub> Receptor Blockers as Potential Antipsychotic Drugs. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2014, 349, 138-154.	2.5	31
25	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
26	Synthesis and biological evaluation of carbon-11 and fluorine-18 labeled tracers for in vivo visualization of PDE10A. <i>Nuclear Medicine and Biology</i> , 2014, 41, 695-704.	0.6	15
27	Discovery of a new series of [1,2,4]triazolo[4,3-a]quinoxalines as dual phosphodiesterase 2/phosphodiesterase 10 (PDE2/PDE10) inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 785-790.	2.2	28
28	Pharmacological Characterization of JNJ-40068782, a New Potent, Selective, and Systemically Active Positive Allosteric Modulator of the mGlu2 Receptor and Its Radioligand [ $^3\text{H}$ ]JNJ-40068782. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2013, 346, 514-527.	2.5	59
29	Differential Interaction of Neuroleptics with Apomorphine-Induced Behavior in Rats as a Function of Changing Levels of Dopamine Receptor Stimulation. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2013, 347, 681-696.	2.5	1
30	Systemic Immune Activation Leads to Neuroinflammation and Sickness Behavior in Mice. <i>Mediators of Inflammation</i> , 2013, 2013, 1-14.	3.0	264
31	Pharmacology of JNJ-37822681, a Specific and Fast-Dissociating D <sub>2</sub> Antagonist for the Treatment of Schizophrenia. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2012, 342, 91-105.	2.5	33
32	Synthesis, Evaluation, and Radiolabeling of New Potent Positive Allosteric Modulators of the Metabotropic Glutamate Receptor 2 as Potential Tracers for Positron Emission Tomography Imaging. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 8685-8699.	6.4	48
33	Synthesis, In Vivo Occupancy, and Radiolabeling of Potent Phosphodiesterase Subtype-10 Inhibitors as Candidates for Positron Emission Tomography Imaging. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 5820-5835.	6.4	43
34	Memantine-induced brain activation as a model for the rapid screening of potential novel antipsychotic compounds: exemplified by activity of an mGlu2/3 receptor agonist. <i>Psychopharmacology</i> , 2011, 214, 505-514.	3.1	32
35	Molecular properties affecting fast dissociation from the D2 receptor. <i>Bioorganic and Medicinal Chemistry</i> , 2011, 19, 2231-2241.	3.0	46
36	Patterns of Brain Glucose Metabolism Induced by Phosphodiesterase 10A Inhibitors in the Mouse: A Potential Translational Biomarker. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2011, 339, 210-217.	2.5	25

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37	Adjunctive $\hat{1}\pm 2$ -adrenoceptor blockade enhances the antipsychotic-like effect of risperidone and facilitates cortical dopaminergic and glutamatergic, NMDA receptor-mediated transmission. <i>International Journal of Neuropsychopharmacology</i> , 2010, 13, 891-903.	2.1	51
38	Preclinical Evaluation of <sup>18</sup> F-JNJ41510417 as a Radioligand for PET Imaging of Phosphodiesterase-10A in the Brain. <i>Journal of Nuclear Medicine</i> , 2010, 51, 1584-1591.	5.0	64
39	In vivo evidence for ligand-specific receptor activation in the central CRF system, as measured by local cerebral glucose utilization. <i>Peptides</i> , 2009, 30, 947-954.	2.4	6
40	Activator of G protein signaling type 3 mRNA is widely distributed in the rat brain and is particularly abundant in the subventricular zone-olfactory bulb system of neural precursor cell proliferation, migration and differentiation. <i>Neuroscience Letters</i> , 2006, 391, 116-121.	2.1	4
41	Combined $\hat{1}\pm 2$ and D2/3 receptor blockade enhances cortical glutamatergic transmission and reverses cognitive impairment in the rat. <i>International Journal of Neuropsychopharmacology</i> , 2005, 8, 315-327.	2.1	70
42	MK-801 alters RGS2 levels and adenylyl cyclase sensitivity in the rat striatum. <i>NeuroReport</i> , 2005, 16, 159-162.	1.2	4
43	Metabotropic glutamate receptor 1 blockade impairs acquisition and retention in a spatial Water maze task. <i>Behavioural Brain Research</i> , 2005, 164, 52-60.	2.2	86
44	Dopamine receptor-mediated regulation of RGS2 and RGS4 mRNA differentially depends on ascending dopamine projections and time. <i>European Journal of Neuroscience</i> , 2004, 19, 2249-2260.	2.6	63
45	Anxiolytic- and antidepressant-like profile of a new CRF1 receptor antagonist, R278995/CRA0450. <i>European Journal of Pharmacology</i> , 2004, 485, 145-158.	3.5	121
46	Metabotropic glutamate 1 receptor distribution and occupancy in the rat brain: a quantitative autoradiographic study using [ <sup>3</sup> H]R214127. <i>Neuropharmacology</i> , 2004, 46, 609-619.	4.1	73
47	JNJ16259685, a highly potent, selective and systemically active mGlu1 receptor antagonist. <i>Neuropharmacology</i> , 2004, 47, 961-972.	4.1	145
48	Striatal gene expression of RGS2 and RGS4 is specifically mediated by dopamine D1 and D2 receptors: clues for RGS2 and RGS4 functions. <i>Journal of Neurochemistry</i> , 2003, 84, 1118-1127.	3.9	95
49	[ <sup>3</sup> H]R214127: A Novel High-Affinity Radioligand for the mGlu1 Receptor Reveals a Common Binding Site Shared by Multiple Allosteric Antagonists. <i>Molecular Pharmacology</i> , 2003, 63, 1082-1093.	2.3	129
50	Differential Membrane Targeting and Pharmacological Characterization of Chimeras of Rat Serotonin 5-HT1A and 5-HT1B Receptors Expressed in Epithelial LLC-PK1 Cells. <i>Journal of Neurochemistry</i> , 2002, 71, 2294-2303.	3.9	16
51	Somatodendritic localization of 5-HT1A and preterminal axonal localization of 5-HT1B serotonin receptors in adult rat brain. <i>Journal of Comparative Neurology</i> , 2000, 417, 181-194.	1.6	402
52	Dominant Role of the Cytosolic C-Terminal Domain of the Rat 5-HT1B Receptor in Axonal Apical Targeting. <i>Journal of Neuroscience</i> , 2000, 20, 9111-9118.	3.6	45
53	Light and electron microscopic immunocytochemical visualization of 5-HT1B receptors in the rat brain. <i>Brain Research</i> , 1997, 760, 281-286.	2.2	70