Jonathan A Javitch

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263 84 22,094 143 h-index g-index citations papers 6.66 24,488 9.7 307 avg, IF L-index ext. citations ext. papers

#	Paper	IF	Citations
263	Parkinsonism-inducing neurotoxin, N-methyl-4-phenyl-1,2,3,6 -tetrahydropyridine: uptake of the metabolite N-methyl-4-phenylpyridine by dopamine neurons explains selective toxicity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1985 , 82, 2173-7	11.5	1024
262	Structure of the human dopamine D3 receptor in complex with a D2/D3 selective antagonist. <i>Science</i> , 2010 , 330, 1091-5	33.3	938
261	Functional selectivity and classical concepts of quantitative pharmacology. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2007 , 320, 1-13	4.7	870
260	Activation of the beta 2-adrenergic receptor involves disruption of an ionic lock between the cytoplasmic ends of transmembrane segments 3 and 6. <i>Journal of Biological Chemistry</i> , 2001 , 276, 2917	1 ⁵ 7 ⁴	490
259	Structural Mimicry in G Protein-Coupled Receptors: Implications of the High-Resolution Structure of Rhodopsin for Structure-Function Analysis of Rhodopsin-Like Receptors. <i>Molecular Pharmacology</i> , 2001 , 60, 1-19	4.3	407
258	Amphetamine-induced loss of human dopamine transporter activity: an internalization-dependent and cocaine-sensitive mechanism. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2000 , 97, 6850-5	11.5	317
257	Building a new conceptual framework for receptor heteromers. <i>Nature Chemical Biology</i> , 2009 , 5, 131-4	11.7	313
256	The mechanism of a neurotransmitter:sodium symporterinward release of Na+ and substrate is triggered by substrate in a second binding site. <i>Molecular Cell</i> , 2008 , 30, 667-77	17.6	308
255	The binding site of aminergic G protein-coupled receptors: the transmembrane segments and second extracellular loop. <i>Annual Review of Pharmacology and Toxicology</i> , 2002 , 42, 437-67	17.9	303
254	Allosteric communication between protomers of dopamine class A GPCR dimers modulates activation. <i>Nature Chemical Biology</i> , 2009 , 5, 688-95	11.7	294
253	Beta2 adrenergic receptor activation. Modulation of the proline kink in transmembrane 6 by a rotamer toggle switch. <i>Journal of Biological Chemistry</i> , 2002 , 277, 40989-96	5.4	288
252	Dopamine D2 receptors form higher order oligomers at physiological expression levels. <i>EMBO Journal</i> , 2008 , 27, 2293-304	13	286
251	Time-resolved FRET between GPCR ligands reveals oligomers in native tissues. <i>Nature Chemical Biology</i> , 2010 , 6, 587-94	11.7	277
250	Discovery of Earrestin-biased dopamine D2 ligands for probing signal transduction pathways essential for antipsychotic efficacy. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 18488-93	11.5	261
249	The binding sites for cocaine and dopamine in the dopamine transporter overlap. <i>Nature Neuroscience</i> , 2008 , 11, 780-9	25.5	260
248	International Union of Basic and Clinical Pharmacology. LXVII. Recommendations for the recognition and nomenclature of G protein-coupled receptor heteromultimers. <i>Pharmacological Reviews</i> , 2007 , 59, 5-13	22.5	255
247	Crosstalk in G protein-coupled receptors: changes at the transmembrane homodimer interface determine activation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005 , 102, 17495-500	11.5	252

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246	Substrate-modulated gating dynamics in a Na+-coupled neurotransmitter transporter homologue. <i>Nature</i> , 2011 , 474, 109-13	50.4	244	
245	The fourth transmembrane segment forms the interface of the dopamine D2 receptor homodimer. Journal of Biological Chemistry, 2003 , 278, 4385-8	5.4	242	
244	Uptake of MPP(+) by dopamine neurons explains selectivity of parkinsonism-inducing neurotoxin, MPTP. <i>European Journal of Pharmacology</i> , 1984 , 106, 455-6	5.3	242	
243	A comprehensive structure-based alignment of prokaryotic and eukaryotic neurotransmitter/Na+symporters (NSS) aids in the use of the LeuT structure to probe NSS structure and function. Molecular Pharmacology, 2006 , 70, 1630-42	4.3	232	
242	Roles of the Akt/GSK-3 and Wnt signaling pathways in schizophrenia and antipsychotic drug action. <i>American Journal of Psychiatry</i> , 2010 , 167, 388-96	11.9	228	
241	Amphetamine induces dopamine efflux through a dopamine transporter channel. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005 , 102, 3495-500	11.5	217	
240	Mitogen-activated protein kinase regulates dopamine transporter surface expression and dopamine transport capacity. <i>Journal of Neuroscience</i> , 2003 , 23, 8480-8	6.6	216	
239	Single-molecule dynamics of gating in a neurotransmitter transporter homologue. <i>Nature</i> , 2010 , 465, 188-93	50.4	213	
238	Treatment resistant depression: A multi-scale, systems biology approach. <i>Neuroscience and Biobehavioral Reviews</i> , 2018 , 84, 272-288	9	209	
237	The role of kinetic context in apparent biased agonism at GPCRs. <i>Nature Communications</i> , 2016 , 7, 108	42 17.4	206	
236	Cyanine fluorophore derivatives with enhanced photostability. <i>Nature Methods</i> , 2011 , 9, 68-71	21.6	203	
235	The second extracellular loop of the dopamine D2 receptor lines the binding-site crevice. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 440-5	11.5	198	
234	Mutation of a highly conserved aspartic acid in the beta2 adrenergic receptor: constitutive activation, structural instability, and conformational rearrangement of transmembrane segment 6. <i>Molecular Pharmacology</i> , 1999 , 56, 175-84	4.3	197	
233	Detection of antigen interactions ex vivo by proximity ligation assay: endogenous dopamine D2-adenosine A2A receptor complexes in the striatum. <i>BioTechniques</i> , 2011 , 51, 111-8	2.5	193	
232	Mechanism of chloride interaction with neurotransmitter:sodium symporters. <i>Nature</i> , 2007 , 449, 726-3	050.4	188	
231	N-terminal phosphorylation of the dopamine transporter is required for amphetamine-induced efflux. <i>PLoS Biology</i> , 2004 , 2, E78	9.7	188	
230	Calmodulin kinase II interacts with the dopamine transporter C terminus to regulate amphetamine-induced reverse transport. <i>Neuron</i> , 2006 , 51, 417-29	13.9	177	
229	Symmetrical dimer of the human dopamine transporter revealed by cross-linking Cys-306 at the extracellular end of the sixth transmembrane segment. <i>Proceedings of the National Academy of Sciences of the United States of America</i> 2001 , 98, 10055, 60	11.5	174	

228	Mapping the binding-site crevice of the dopamine D2 receptor by the substituted-cysteine accessibility method. <i>Neuron</i> , 1995 , 14, 825-31	13.9	170
227	PI 3-kinase regulation of dopamine uptake. <i>Journal of Neurochemistry</i> , 2002 , 81, 859-69	6	168
226	Single-molecule analysis of ligand efficacy in AR-G-protein activation. <i>Nature</i> , 2017 , 547, 68-73	50.4	164
225	Constitutive activation of the beta2 adrenergic receptor alters the orientation of its sixth membrane-spanning segment. <i>Journal of Biological Chemistry</i> , 1997 , 272, 18546-9	5.4	163
224	Synthetic and Receptor Signaling Explorations of the Mitragyna Alkaloids: Mitragynine as an Atypical Molecular Framework for Opioid Receptor Modulators. <i>Journal of the American Chemical Society</i> , 2016 , 138, 6754-64	16.4	161
223	Binding of an octylglucoside detergent molecule in the second substrate (S2) site of LeuT establishes an inhibitor-bound conformation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009 , 106, 5563-8	11.5	160
222	The organic cation transporter-3 is a pivotal modulator of neurodegeneration in the nigrostriatal dopaminergic pathway. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009 , 106, 8043-8	11.5	160
221	A cluster of aromatic residues in the sixth membrane-spanning segment of the dopamine D2 receptor is accessible in the binding-site crevice. <i>Biochemistry</i> , 1998 , 37, 998-1006	3.2	160
220	Reaction of oxidized dopamine with endogenous cysteine residues in the human dopamine transporter. <i>Journal of Neurochemistry</i> , 2001 , 76, 1242-51	6	158
219	Ion/substrate-dependent conformational dynamics of a bacterial homolog of neurotransmitter:sodium symporters. <i>Nature Structural and Molecular Biology</i> , 2010 , 17, 822-9	17.6	157
218	D2 receptors regulate dopamine transporter function via an extracellular signal-regulated kinases 1 and 2-dependent and phosphoinositide 3 kinase-independent mechanism. <i>Molecular Pharmacology</i> , 2007 , 71, 1222-32	4.3	153
217	Amphetamine-induced dopamine efflux. A voltage-sensitive and intracellular Na+-dependent mechanism. <i>Journal of Biological Chemistry</i> , 2003 , 278, 12070-7	5.4	152
216	The forgotten serine. A critical role for Ser-2035.42 in ligand binding to and activation of the beta 2-adrenergic receptor. <i>Journal of Biological Chemistry</i> , 2000 , 275, 37779-88	5.4	147
215	Cocaine increases dopamine uptake and cell surface expression of dopamine transporters. <i>Biochemical and Biophysical Research Communications</i> , 2002 , 290, 1545-50	3.4	146
214	Flotillin-1 is essential for PKC-triggered endocytosis and membrane microdomain localization of DAT. <i>Nature Neuroscience</i> , 2011 , 14, 469-77	25.5	141
213	Monitoring the function of membrane transport proteins in detergent-solubilized form. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007 , 104, 3603-8	11.5	140
212	N-terminal truncation of the dopamine transporter abolishes phorbol ester- and substance P receptor-stimulated phosphorylation without impairing transporter internalization. <i>Journal of Biological Chemistry</i> , 2003 , 278, 4990-5000	5.4	138
211	Increasing dopamine D2 receptor expression in the adult nucleus accumbens enhances motivation. <i>Molecular Psychiatry</i> , 2013 , 18, 1025-33	15.1	137

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210	Amphetamine and methamphetamine differentially affect dopamine transporters in vitro and in vivo. <i>Journal of Biological Chemistry</i> , 2009 , 284, 2978-2989	5.4	135
209	Molecular determinants of selectivity and efficacy at the dopamine D3 receptor. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 6689-99	8.3	131
208	Signaling pathways in schizophrenia: emerging targets and therapeutic strategies. <i>Trends in Pharmacological Sciences</i> , 2010 , 31, 381-90	13.2	128
207	Hetero-oligomerization of CCR2, CCR5, and CXCR4 and the protean effects of "selective" antagonists. <i>Journal of Biological Chemistry</i> , 2009 , 284, 31270-9	5.4	128
206	A cysteine residue in the third membrane-spanning segment of the human D2 dopamine receptor is exposed in the binding-site crevice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1994 , 91, 10355-9	11.5	127
205	Paraquat neurotoxicity is mediated by the dopamine transporter and organic cation transporter-3. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 20766-71	11.5	126
204	The Behavioral Effects of the Antidepressant Tianeptine Require the Mu-Opioid Receptor. <i>Neuropsychopharmacology</i> , 2017 , 42, 2052-2063	8.7	123
203	Akt is essential for insulin modulation of amphetamine-induced human dopamine transporter cell-surface redistribution. <i>Molecular Pharmacology</i> , 2005 , 68, 102-9	4.3	121
202	A mechanism for intracellular release of Na+ by neurotransmitter/sodium symporters. <i>Nature Structural and Molecular Biology</i> , 2014 , 21, 1006-12	17.6	119
201	Cocaine alters the accessibility of endogenous cysteines in putative extracellular and intracellular loops of the human dopamine transporter. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1998 , 95, 9238-43	11.5	118
200	Residues in the seventh membrane-spanning segment of the dopamine D2 receptor accessible in the binding-site crevice. <i>Biochemistry</i> , 1996 , 35, 11278-85	3.2	116
199	Structural mimicry in G protein-coupled receptors: implications of the high-resolution structure of rhodopsin for structure-function analysis of rhodopsin-like receptors. <i>Molecular Pharmacology</i> , 2001 , 60, 1-19	4.3	114
198	Sodium-dependent neurotransmitter transporters: oligomerization as a determinant of transporter function and trafficking. <i>Molecular Interventions: Pharmacological Perspectives From Biology, Chemistry and Genomics,</i> 2004 , 4, 38-47		110
197	An intracellular interaction network regulates conformational transitions in the dopamine transporter. <i>Journal of Biological Chemistry</i> , 2008 , 283, 17691-701	5.4	108
196	Conformational dynamics of ligand-dependent alternating access in LeuT. <i>Nature Structural and Molecular Biology</i> , 2014 , 21, 472-9	17.6	102
195	Residues in the fifth membrane-spanning segment of the dopamine D2 receptor exposed in the binding-site crevice. <i>Biochemistry</i> , 1995 , 34, 16433-9	3.2	102
194	Crystal structure of a potassium ion transporter, TrkH. <i>Nature</i> , 2011 , 471, 336-40	50.4	101
193	The substrate-driven transition to an inward-facing conformation in the functional mechanism of the dopamine transporter. <i>PLoS ONE</i> , 2011 , 6, e16350	3.7	99

192	Human immunodeficiency virus (HIV) infection of human macrophages is increased by dopamine: a bridge between HIV-associated neurologic disorders and drug abuse. <i>American Journal of Pathology</i> , 2009 , 175, 1148-59	5.8	98
191	Syntaxin 1A interaction with the dopamine transporter promotes amphetamine-induced dopamine efflux. <i>Molecular Pharmacology</i> , 2008 , 74, 1101-8	4.3	98
190	The human dopamine transporter forms a tetramer in the plasma membrane: cross-linking of a cysteine in the fourth transmembrane segment is sensitive to cocaine analogs. <i>Journal of Biological Chemistry</i> , 2003 , 278, 45045-8	5.4	98
189	Extrapyramidal side effects of antipsychotics are linked to their association kinetics at dopamine D receptors. <i>Nature Communications</i> , 2017 , 8, 763	17.4	97
188	Identification of novel functionally selective Eppioid receptor scaffolds. <i>Molecular Pharmacology</i> , 2014 , 85, 83-90	4.3	95
187	A new mechanism of allostery in a G protein-coupled receptor dimer. <i>Nature Chemical Biology</i> , 2014 , 10, 745-52	11.7	95
186	CODA-RET reveals functional selectivity as a result of GPCR heteromerization. <i>Nature Chemical Biology</i> , 2011 , 7, 624-30	11.7	92
185	Peroxynitrite inactivates the human dopamine transporter by modification of cysteine 342: potential mechanism of neurotoxicity in dopamine neurons. <i>Journal of Neuroscience</i> , 2002 , 22, 4399-405	-6.6 -	91
184	Dysregulation of dopamine transporters via dopamine D2 autoreceptors triggers anomalous dopamine efflux associated with attention-deficit hyperactivity disorder. <i>Journal of Neuroscience</i> , 2010 , 30, 6048-57	6.6	90
183	The uptake inhibitors cocaine and benztropine differentially alter the conformation of the human dopamine transporter. <i>Journal of Biological Chemistry</i> , 2001 , 276, 29012-8	5.4	88
182	Dopamine D4/D2 receptor selectivity is determined by A divergent aromatic microdomain contained within the second, third, and seventh membrane-spanning segments. <i>Molecular Pharmacology</i> , 1999 , 56, 1116-26	4.3	88
181	The ants go marching two by two: oligomeric structure of G-protein-coupled receptors. <i>Molecular Pharmacology</i> , 2004 , 66, 1077-82	4.3	85
180	[3H]mazindol binding associated with neuronal dopamine uptake sites in corpus striatum membranes. <i>European Journal of Pharmacology</i> , 1983 , 90, 461-2	5.3	84
179	PIP2 regulates psychostimulant behaviors through its interaction with a membrane protein. <i>Nature Chemical Biology</i> , 2014 , 10, 582-589	11.7	83
178	Evidence against dopamine D1/D2 receptor heteromers. <i>Molecular Psychiatry</i> , 2015 , 20, 1373-85	15.1	83
177	Experimental conditions can obscure the second high-affinity site in LeuT. <i>Nature Structural and Molecular Biology</i> , 2012 , 19, 207-11	17.6	82
176	6PGuanidinonaltrindole (6PGNTI) is a G protein-biased Eppioid receptor agonist that inhibits arrestin recruitment. <i>Journal of Biological Chemistry</i> , 2012 , 287, 27050-4	5.4	82
175	Antipsychotic drug mechanisms: links between therapeutic effects, metabolic side effects and the insulin signaling pathway. <i>Molecular Psychiatry</i> , 2008 , 13, 918-29	15.1	82

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174	Intracellular Ca2+ regulates amphetamine-induced dopamine efflux and currents mediated by the human dopamine transporter. <i>Molecular Pharmacology</i> , 2004 , 66, 137-43	4.3	81	
173	Identification of intracellular residues in the dopamine transporter critical for regulation of transporter conformation and cocaine binding. <i>Journal of Biological Chemistry</i> , 2004 , 279, 3228-38	5.4	79	
172	Regulation of dopamine transporter function and cell surface expression by D3 dopamine receptors. <i>Journal of Biological Chemistry</i> , 2007 , 282, 35842-54	5.4	78	
171	G protein-coupled receptor kinase-mediated phosphorylation regulates post-endocytic trafficking of the D2 dopamine receptor. <i>Journal of Biological Chemistry</i> , 2009 , 284, 15038-51	5.4	77	
170	State-dependent conformations of the translocation pathway in the tyrosine transporter Tyt1, a novel neurotransmitter:sodium symporter from Fusobacterium nucleatum. <i>Journal of Biological Chemistry</i> , 2006 , 281, 26444-54	5.4	77	
169	Synergistic contributions of the functional groups of epinephrine to its affinity and efficacy at the beta2 adrenergic receptor. <i>Molecular Pharmacology</i> , 2004 , 65, 1181-90	4.3	77	
168	Amphetamine regulation of dopamine transport. Combined measurements of transporter currents and transporter imaging support the endocytosis of an active carrier. <i>Journal of Biological Chemistry</i> , 2004 , 279, 8966-75	5.4	77	
167	Characterization of a functional bacterial homologue of sodium-dependent neurotransmitter transporters. <i>Journal of Biological Chemistry</i> , 2003 , 278, 12703-9	5.4	77	
166	Dual agonist occupancy of AT1-R-IZC-AR heterodimers results in atypical Gs-PKA signaling. <i>Nature Chemical Biology</i> , 2015 , 11, 271-9	11.7	76	
165	7-Hydroxymitragynine Is an Active Metabolite of Mitragynine and a Key Mediator of Its Analgesic Effects. <i>ACS Central Science</i> , 2019 , 5, 992-1001	16.8	75	
164	What can crystal structures of aminergic receptors tell us about designing subtype-selective ligands?. <i>Pharmacological Reviews</i> , 2015 , 67, 198-213	22.5	75	
163	The atypical antidepressant and neurorestorative agent tianeptine is a Eppioid receptor agonist. <i>Translational Psychiatry</i> , 2014 , 4, e411	8.6	74	
162	Surface targeting of the dopamine transporter involves discrete epitopes in the distal C terminus but does not require canonical PDZ domain interactions. <i>Journal of Neuroscience</i> , 2004 , 24, 7024-36	6.6	72	
161	GPCR-OKB: the G Protein Coupled Receptor Oligomer Knowledge Base. <i>Bioinformatics</i> , 2010 , 26, 1804-5	57.2	71	
160	Mechanisms of amphetamine action illuminated through optical monitoring of dopamine synaptic vesicles in Drosophila brain. <i>Nature Communications</i> , 2016 , 7, 10652	17.4	70	
159	Electrostatic and aromatic microdomains within the binding-site crevice of the D2 receptor: contributions of the second membrane-spanning segment. <i>Biochemistry</i> , 1999 , 38, 7961-8	3.2	68	
158	Crystal structure of a phosphorylation-coupled saccharide transporter. <i>Nature</i> , 2011 , 473, 50-4	50.4	67	
157	Making structural sense of dimerization interfaces of delta opioid receptor homodimers. <i>Biochemistry</i> , 2011 , 50, 1682-90	3.2	66	

156	Chloride binding site of neurotransmitter sodium symporters. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013 , 110, 8489-94	11.5	65
155	The fourth transmembrane segment of the dopamine D2 receptor: accessibility in the binding-site crevice and position in the transmembrane bundle. <i>Biochemistry</i> , 2000 , 39, 12190-9	3.2	64
154	CrossTalk opposing view: Weighing the evidence for class A GPCR dimers, the jury is still out. <i>Journal of Physiology</i> , 2014 , 592, 2443-5	3.9	63
153	Impact of D2 receptor internalization on binding affinity of neuroimaging radiotracers. <i>Neuropsychopharmacology</i> , 2010 , 35, 806-17	8.7	63
152	Dopamine-mediated autocrine inhibitory circuit regulating human insulin secretion in vitro. <i>Molecular Endocrinology</i> , 2012 , 26, 1757-72		61
151	Recruitment of beta-arrestin2 to the dopamine D2 receptor: insights into anti-psychotic and anti-parkinsonian drug receptor signaling. <i>Neuropharmacology</i> , 2008 , 54, 1215-22	5.5	61
150	Discovery and characterization of a G protein-biased agonist that inhibits Earrestin recruitment to the D2 dopamine receptor. <i>Molecular Pharmacology</i> , 2014 , 86, 96-105	4.3	59
149	G protein-coupled receptor kinase-2 constitutively regulates D2 dopamine receptor expression and signaling independently of receptor phosphorylation. <i>Journal of Biological Chemistry</i> , 2009 , 284, 34103-	.15 ⁴	58
148	Regulation of dopamine transporter trafficking by intracellular amphetamine. <i>Molecular Pharmacology</i> , 2006 , 70, 542-8	4.3	57
147	A juxtamembrane mutation in the N terminus of the dopamine transporter induces preference for an inward-facing conformation. <i>Molecular Pharmacology</i> , 2009 , 75, 514-24	4.3	56
146	The first transmembrane segment of the dopamine D2 receptor: accessibility in the binding-site crevice and position in the transmembrane bundle. <i>Biochemistry</i> , 2001 , 40, 12339-48	3.2	56
145	Mechanisms of inverse agonism of antipsychotic drugs at the D(2) dopamine receptor: use of a mutant D(2) dopamine receptor that adopts the activated conformation. <i>Journal of Neurochemistry</i> , 2001 , 77, 493-504	6	55
144	Discovery of a novel selective kappa-opioid receptor agonist using crystal structure-based virtual screening. <i>Journal of Chemical Information and Modeling</i> , 2013 , 53, 521-6	6.1	54
143	Currents in response to rapid concentration jumps of amphetamine uncover novel aspects of human dopamine transporter function. <i>Journal of Neuroscience</i> , 2008 , 28, 976-89	6.6	54
142	Structure and functional interaction of the extracellular domain of human GABA(B) receptor GBR2. <i>Nature Neuroscience</i> , 2012 , 15, 970-8	25.5	53
141	Imaging the high-affinity state of the dopamine D2 receptor in vivo: fact or fiction?. <i>Biochemical Pharmacology</i> , 2012 , 83, 193-8	6	51
140	Transport-dependent accessibility of a cytoplasmic loop cysteine in the human dopamine transporter. <i>Journal of Biological Chemistry</i> , 2000 , 275, 1608-14	5.4	51
139	The tetrahydroisoquinoline derivative SB269,652 is an allosteric antagonist at dopamine D3 and D2 receptors. <i>Molecular Pharmacology</i> , 2010 , 78, 925-34	4.3	49

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138	Accumbens dopamine D2 receptors increase motivation by decreasing inhibitory transmission to the ventral pallidum. <i>Nature Communications</i> , 2018 , 9, 1086	17.4	48
137	Substrate-dependent proton antiport in neurotransmitter:sodium symporters. <i>Nature Chemical Biology</i> , 2010 , 6, 109-16	11.7	48
136	Use of the substituted cysteine accessibility method to study the structure and function of G protein-coupled receptors. <i>Methods in Enzymology</i> , 2002 , 343, 137-56	1.7	48
135	Development and Antiparkinsonian Activity of VU0418506, a Selective Positive Allosteric Modulator of Metabotropic Glutamate Receptor 4 Homomers without Activity at mGlu2/4 Heteromers. <i>ACS Chemical Neuroscience</i> , 2016 , 7, 1201-11	5.7	47
134	Presynaptic regulation of dopamine transmission in schizophrenia. Schizophrenia Bulletin, 2011, 37, 108	-173	47
133	Probing structure of neurotransmitter transporters by substituted-cysteine accessibility method. <i>Methods in Enzymology</i> , 1998 , 296, 331-46	1.7	46
132	A single glycine in extracellular loop 1 is the critical determinant for pharmacological specificity of dopamine D2 and D3 receptors. <i>Molecular Pharmacology</i> , 2013 , 84, 854-64	4.3	45
131	Cholinergic agonists as novel treatments for schizophrenia: the promise of rational drug development for psychiatry. <i>American Journal of Psychiatry</i> , 2008 , 165, 931-6	11.9	45
130	Dopamine receptor activation increases HIV entry into primary human macrophages. <i>PLoS ONE</i> , 2014 , 9, e108232	3.7	45
129	Optical Control of Dopamine Receptors Using a Photoswitchable Tethered Inverse Agonist. <i>Journal of the American Chemical Society</i> , 2017 , 139, 18522-18535	16.4	43
128	Neuronal Depolarization Drives Increased Dopamine Synaptic Vesicle Loading via VGLUT. <i>Neuron</i> , 2017 , 95, 1074-1088.e7	13.9	42
127	Mechanism of the Association between Na+ Binding and Conformations at the Intracellular Gate in Neurotransmitter:Sodium Symporters. <i>Journal of Biological Chemistry</i> , 2015 , 290, 13992-4003	5.4	41
126	Regional Heterogeneity of D2-Receptor Signaling in the Dorsal Striatum and Nucleus Accumbens. <i>Neuron</i> , 2018 , 98, 575-587.e4	13.9	41
125	How did the neurotransmitter cross the bilayer? A closer view. <i>Current Opinion in Neurobiology</i> , 2005 , 15, 296-304	7.6	37
124	Metabotropic Glutamate Receptor 5 and Glutamate Involvement in Major Depressive Disorder: A Multimodal Imaging Study. <i>Biological Psychiatry: Cognitive Neuroscience and Neuroimaging</i> , 2017 , 2, 449-	-45ts	36
123	Electronic tuning of self-healing fluorophores for live-cell and single-molecule imaging. <i>Chemical Science</i> , 2017 , 8, 755-762	9.4	36
122	Structure-activity relationships for a novel series of citalopram (1-(3-(dimethylamino)propyl)-1-(4-fluorophenyl)-1,3-dihydroisobenzofuran-5-carbonitrile) analogues at monoamine transporters. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 6112-21	8.3	36
121	A pincer-like configuration of TM2 in the human dopamine transporter is responsible for indirect effects on cocaine binding. <i>Neuropharmacology</i> , 2005 , 49, 780-90	5.5	36

120	Probing conformational changes in neurotransmitter transporters: a structural context. <i>European Journal of Pharmacology</i> , 2003 , 479, 3-12	5.3	36
119	Potentiating SLC transporter activity: Emerging drug discovery opportunities. <i>Biochemical Pharmacology</i> , 2017 , 135, 1-11	6	35
118	High Affinity Dopamine D3 Receptor (D3R)-Selective Antagonists Attenuate Heroin Self-Administration in Wild-Type but not D3R Knockout Mice. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 6195-213	8.3	35
117	Upregulation of dopamine D2 receptors in the nucleus accumbens indirect pathway increases locomotion but does not reduce alcohol consumption. <i>Neuropsychopharmacology</i> , 2015 , 40, 1609-18	8.7	34
116	In vivo variation in same-day estimates of metabotropic glutamate receptor subtype 5 binding using [C] ABP688 and [F] FPEB. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2017 , 37, 2716-2727	7.3	34
115	PI3K signaling supports amphetamine-induced dopamine efflux. <i>Biochemical and Biophysical Research Communications</i> , 2008 , 372, 656-61	3.4	34
114	Toward Understanding the Structural Basis of Partial Agonism at the Dopamine D Receptor. Journal of Medicinal Chemistry, 2017 , 60, 580-593	8.3	33
113	Genetically Targeted Optical Control of an Endogenous G Protein-Coupled Receptor. <i>Journal of the American Chemical Society</i> , 2019 , 141, 11522-11530	16.4	32
112	Structure of human GABA receptor in an inactive state. <i>Nature</i> , 2020 , 584, 304-309	50.4	32
111	Using Bioluminescence Resonance Energy Transfer (BRET) to Characterize Agonist-Induced Arrestin Recruitment to Modified and Unmodified G Protein-Coupled Receptors. <i>Current Protocols in Pharmacology</i> , 2015 , 70, 2.14.1-2.14.14	4.1	32
110	The conserved cysteine 7.38 residue is differentially accessible in the binding-site crevices of the mu, delta, and kappa opioid receptors. <i>Biochemistry</i> , 2000 , 39, 13904-15	3.2	32
109	Arrestin recruitment to dopamine D2 receptor mediates locomotion but not incentive motivation. <i>Molecular Psychiatry</i> , 2020 , 25, 2086-2100	15.1	32
108	Zn(2+) site engineering at the oligomeric interface of the dopamine transporter. <i>FEBS Letters</i> , 2002 , 524, 87-91	3.8	31
107	Comparison of the amino acid residues in the sixth transmembrane domains accessible in the binding-site crevices of mu, delta, and kappa opioid receptors. <i>Biochemistry</i> , 2001 , 40, 8018-29	3.2	30
106	Chloroethylclonidine and 2-aminoethyl methanethiosulfonate recognize two different conformations of the human alpha(2A)-adrenergic receptor. <i>Journal of Biological Chemistry</i> , 1999 , 274, 21867-72	5.4	30
105	Single-molecule FRET imaging of GPCR dimers in living cells. <i>Nature Methods</i> , 2021 , 18, 397-405	21.6	30
104	New roles for dopamine D and D receptors in pancreatic beta cell insulin secretion. <i>Molecular Psychiatry</i> , 2020 , 25, 2070-2085	15.1	30
103	The membrane protein LeuT in micellar systems: aggregation dynamics and detergent binding to the S2 site. <i>Journal of the American Chemical Society</i> , 2013 , 135, 14266-75	16.4	29

102	Substrate-induced unlocking of the inner gate determines the catalytic efficiency of a neurotransmitter:sodium symporter. <i>Journal of Biological Chemistry</i> , 2015 , 290, 26725-38	5.4	28	
101	A partially-open inward-facing intermediate conformation of LeuT is associated with Na release and substrate transport. <i>Nature Communications</i> , 2018 , 9, 230	17.4	28	
100	Yohimbine depresses excitatory transmission in BNST and impairs extinction of cocaine place preference through orexin-dependent, norepinephrine-independent processes. Neuropsychopharmacology, 2012, 37, 2253-66	8.7	28	
99	Cannabinoid CB1 and CB2 Receptor-Mediated Arrestin Translocation: Species, Subtype, and Agonist-Dependence. <i>Frontiers in Pharmacology</i> , 2019 , 10, 350	5.6	26	
98	Novel Analogues of (R)-5-(Methylamino)-5,6-dihydro-4H-imidazo[4,5,1-ij]quinolin-2(1H)-one (Sumanirole) Provide Clues to Dopamine D2/D3 Receptor Agonist Selectivity. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 2973-88	8.3	26	
97	Segregation of family A G protein-coupled receptor protomers in the plasma membrane. <i>Molecular Pharmacology</i> , 2013 , 84, 346-52	4.3	25	
96	Gs-Iversus Golf-dependent functional selectivity mediated by the dopamine D receptor. <i>Nature Communications</i> , 2018 , 9, 486	17.4	24	
95	Mutation of three residues in the third intracellular loop of the dopamine D2 receptor creates an internalization-defective receptor. <i>Journal of Biological Chemistry</i> , 2014 , 289, 33663-75	5.4	24	
94	Akt-dependent and isoform-specific regulation of dopamine transporter cell surface expression. <i>ACS Chemical Neuroscience</i> , 2010 , 1, 476-81	5.7	24	
93	Dopamine D2 Receptors in the Paraventricular Thalamus Attenuate Cocaine Locomotor Sensitization. <i>ENeuro</i> , 2017 , 4,	3.9	22	
92	Quantifying secondary transport at single-molecule resolution. <i>Nature</i> , 2019 , 575, 528-534	50.4	22	
91	Do Toxic Synthetic Cannabinoid Receptor Agonists Have Signature in Vitro Activity Profiles? A Case Study of AMB-FUBINACA. <i>ACS Chemical Neuroscience</i> , 2019 , 10, 4350-4360	5.7	21	
90	Evidence for limited D1 and D2 receptor coexpression and colocalization within the dorsal striatum of the neonatal mouse. <i>Journal of Comparative Neurology</i> , 2015 , 523, 1175-89	3.4	21	
89	Two allelic isoforms of the serotonin transporter from Schistosoma mansoni display electrogenic transport and high selectivity for serotonin. <i>European Journal of Pharmacology</i> , 2009 , 616, 48-57	5.3	21	
88	Dopamine prevents nitration of tyrosine hydroxylase by peroxynitrite and nitrogen dioxide: is nitrotyrosine formation an early step in dopamine neuronal damage?. <i>Journal of Biological Chemistry</i> , 2003 , 278, 28736-42	5.4	21	
87	Ribosome-associated vesicles: A dynamic subcompartment of the endoplasmic reticulum in secretory cells. <i>Science Advances</i> , 2020 , 6, eaay9572	14.3	20	
86	Characterization of in vivo pharmacokinetic properties of the dopamine D1 receptor agonist DAR-0100A in nonhuman primates using PET with [11C] NNC112 and [11C] raclopride. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2011 , 31, 293-304	7.3	19	
85	The structural determinants of the bitopic binding mode of a negative allosteric modulator of the dopamine D receptor. <i>Biochemical Pharmacology</i> , 2018 , 148, 315-328	6	18	

84	Development of a Rapid Insulin Assay by Homogenous Time-Resolved Fluorescence. <i>PLoS ONE</i> , 2016 , 11, e0148684	3.7	18
83	Distinct inactive conformations of the dopamine D2 and D3 receptors correspond to different extents of inverse agonism. <i>ELife</i> , 2020 , 9,	8.9	18
82	The role of transmembrane segment 5 (TM5) in Na2 release and the conformational transition of neurotransmitter:sodium symporters toward the inward-open state. <i>Journal of Biological Chemistry</i> , 2017 , 292, 7372-7384	5.4	17
81	Tuning the Baird aromatic triplet-state energy of cyclooctatetraene to maximize the self-healing mechanism in organic fluorophores. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020 , 117, 24305-24315	11.5	17
80	Dopamine D2 receptor overexpression in the nucleus accumbens core induces robust weight loss during scheduled fasting selectively in female mice. <i>Molecular Psychiatry</i> , 2021 , 26, 3765-3777	15.1	17
79	Conformational Dynamics on the Extracellular Side of LeuT Controlled by Na+ and K+ Ions and the Protonation State of Glu290. <i>Journal of Biological Chemistry</i> , 2016 , 291, 19786-99	5.4	16
78	The allosteric mechanism of substrate-specific transport in SLC6 is mediated by a volumetric sensor. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 1594	47-1· 5 9:	56 ¹⁶
77	Controlling opioid receptor functional selectivity by targeting distinct subpockets of the orthosteric site. <i>ELife</i> , 2021 , 10,	8.9	16
76	X-ray structure of LeuT in an inward-facing occluded conformation reveals mechanism of substrate release. <i>Nature Communications</i> , 2020 , 11, 1005	17.4	15
75	Signalling profiles of a structurally diverse panel of synthetic cannabinoid receptor agonists. <i>Biochemical Pharmacology</i> , 2020 , 175, 113871	6	15
74	Differential visualization of dopamine D2 and D3 receptors in rat brain. <i>European Journal of Pharmacology</i> , 1993 , 234, 269-72	5.3	15
73	The E2.65A mutation disrupts dynamic binding poses of SB269652 at the dopamine D2 and D3 receptors. <i>PLoS Computational Biology</i> , 2018 , 14, e1005948	5	15
72	The LeuT-fold neurotransmitter:sodium symporter MhsT has two substrate sites. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018 , 115, E7924-E7931	11.5	14
71	Molecular Determinants of the Intrinsic Efficacy of the Antipsychotic Aripiprazole. <i>ACS Chemical Biology</i> , 2019 , 14, 1780-1792	4.9	14
70	The action of a negative allosteric modulator at the dopamine D receptor is dependent upon sodium ions. <i>Scientific Reports</i> , 2018 , 8, 1208	4.9	13
69	Mapping the binding-site crevice of the D2 receptor. <i>Advances in Pharmacology</i> , 1998 , 42, 412-5	5.7	13
68	Agonist-induced formation of unproductive receptor-G complexes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020 , 117, 21723-21730	11.5	13
67	Come Fly with Me: An overview of dopamine receptors in Drosophila melanogaster. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2020 , 126 Suppl 6, 56-65	3.1	13

66	Rebuttal from Nevin A. Lambert and Jonathan A. Javitch. <i>Journal of Physiology</i> , 2014 , 592, 2449	3.9	12	
65	G12/13 is activated by acute tethered agonist exposure in the adhesion GPCR ADGRL3. <i>Nature Chemical Biology</i> , 2020 , 16, 1343-1350	11.7	12	
64	Intramolecular cross-linking in a bacterial homolog of mammalian SLC6 neurotransmitter transporters suggests an evolutionary conserved role of transmembrane segments 7 and 8. <i>Neuropharmacology</i> , 2005 , 49, 715-23	5.5	11	
63	Role of Tau Protein in Remodeling of Circadian Neuronal Circuits and Sleep. <i>Frontiers in Aging Neuroscience</i> , 2019 , 11, 320	5.3	11	
62	Role of Annular Lipids in the Functional Properties of Leucine Transporter LeuT Proteomicelles. <i>Biochemistry</i> , 2016 , 55, 850-9	3.2	10	
61	Luciferase complementation based-detection of G-protein-coupled receptor activity. <i>BioTechniques</i> , 2018 , 65, 9-14	2.5	10	
60	Conformational changes in dopamine transporter intracellular regions upon cocaine binding and dopamine translocation. <i>Neurochemistry International</i> , 2014 , 73, 4-15	4.4	10	
59	[3H]MFZ 2-12: a novel radioligand for the dopamine transporter. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2001 , 11, 1659-61	2.9	10	
58	A Novel Mitragynine Analog with Low-Efficacy Mu Opioid Receptor Agonism Displays Antinociception with Attenuated Adverse Effects. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 13873-1389	92 ^{8.3}	10	
57	The differential actions of clozapine and other antipsychotic drugs on the translocation of dopamine D2 receptors to the cell surface. <i>Journal of Biological Chemistry</i> , 2019 , 294, 5604-5615	5.4	9	
56	Site selective C-H functionalization of Mitragyna alkaloids reveals a molecular switch for tuning opioid receptor signaling efficacy. <i>Nature Communications</i> , 2021 , 12, 3858	17.4	9	
55	Development of novel biosensors to study receptor-mediated activation of the G-protein I subunits G and G. <i>Journal of Biological Chemistry</i> , 2017 , 292, 19989-19998	5.4	8	
54	The substituted-cysteine accessibility method (SCAM) to elucidate membrane protein structure. <i>Current Protocols in Neuroscience</i> , 2001 , Chapter 4, Unit 4.15	2.7	8	
53	Exploring the binding site crevice of a family B G protein-coupled receptor, the type 1 corticotropin releasing factor receptor. <i>Molecular Pharmacology</i> , 2010 , 78, 785-93	4.3	7	
52	Phosphorylation of the Amino Terminus of the Dopamine Transporter: Regulatory Mechanisms and Implications for Amphetamine Action. <i>Advances in Pharmacology</i> , 2018 , 82, 205-234	5.7	6	
51	A role for information collection, management, and integration in structure-function studies of G-protein coupled receptors. <i>Current Pharmaceutical Design</i> , 2006 , 12, 1771-83	3.3	6	
50	Input-specific regulation of glutamatergic synaptic transmission in the medial prefrontal cortex by mGlu/mGlu receptor heterodimers. <i>Science Signaling</i> , 2021 , 14,	8.8	5	
49	New phosphosite-specific antibodies to unravel the role of GRK phosphorylation in dopamine D receptor regulation and signaling. <i>Scientific Reports</i> , 2021 , 11, 8288	4.9	5	

48	Cortical overgrowth in a preclinical forebrain organoid model of CNTNAP2-associated autism spectrum disorder. <i>Nature Communications</i> , 2021 , 12, 4087	17.4	5
47	A novel luminescence-based Earrestin recruitment assay for unmodified receptors. <i>Journal of Biological Chemistry</i> , 2021 , 296, 100503	5.4	5
46	Phospho-specific antibodies targeting the amino terminus of the human dopamine transporter. Journal of Chemical Neuroanatomy, 2017 , 83-84, 91-98	3.2	4
45	Oligomerization Domains of G Protein-Coupled Receptors 2005 , 243-265		4
44	A non-helical region in transmembrane helix 6 of hydrophobic amino acid transporter MhsT mediates substrate recognition. <i>EMBO Journal</i> , 2021 , 40, e105164	13	4
43	Detecting G protein-coupled receptor complexes in postmortem human brain with proximity ligation assay and a Bayesian classifier. <i>BioTechniques</i> , 2020 , 68, 122-129	2.5	4
42	Measuring the effects of ketamine on mGluR5 using [F]FPEB and PET. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2020 , 40, 2254-2264	7.3	4
41	Detection of G Protein-Coupled Receptor Complexes in Postmortem Human Brain by Proximity Ligation Assay. <i>Current Protocols in Neuroscience</i> , 2020 , 91, e86	2.7	3
40	Drs. Lieberman, Javitch, and Moore Reply. American Journal of Psychiatry, 2009, 166, 111-113	11.9	3
39	Dopamine D2 receptors modulate the cholinergic pause and inhibitory learning. <i>Molecular Psychiatry</i> , 2021 ,	15.1	3
38	Disrupting D1-NMDA or D2-NMDA receptor heteromerization prevents cocaine® rewarding effects but preserves natural reward processing. <i>Science Advances</i> , 2021 , 7, eabg5970	14.3	3
37	Structural Basis of Dopamine Receptor Activation 2010 , 47-73		3
36	Synthesis and pharmacological evaluation of bivalent tethered ligands to target the mGlu heterodimeric receptor results in a compound with mGlu homodimer selectivity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020 , 30, 127212	2.9	3
35	Novel Fluorescent Ligands Enable Single-Molecule Localization Microscopy of the Dopamine Transporter. <i>ACS Chemical Neuroscience</i> , 2020 , 11, 3288-3300	5.7	3
34	Crystal structures of LeuT reveal conformational dynamics in the outward-facing states. <i>Journal of Biological Chemistry</i> , 2021 , 296, 100609	5.4	3
33	Exploring Substrate Binding in the Extracellular Vestibule of MhsT by Atomistic Simulations and Markov Models. <i>Journal of Chemical Information and Modeling</i> , 2018 , 58, 1244-1252	6.1	3
32	Mu opioid receptors on hippocampal GABAergic interneurons are critical for the antidepressant effects of tianeptine. <i>Neuropsychopharmacology</i> , 2021 ,	8.7	3
31	Extreme Vetting of Dopamine Receptor Oligomerization 2017 , 99-127		2

(2011-2021)

30	Delineating the interactions between the cannabinoid CB receptor and its regulatory effectors; Earrestins and G protein-coupled receptor kinases. <i>British Journal of Pharmacology</i> , 2021 ,	8.6	2
29	Dopamine D2 receptors modulate the cholinergic pause and inhibitory learning		2
28	How changes in dopamine D2 receptor levels alter striatal circuit function and motivation. <i>Molecular Psychiatry</i> , 2021 ,	15.1	2
27	Sensing conformational changes in metabotropic glutamate receptors. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013 , 110, 5742-3	11.5	1
26	TRAC: A Platform for Structure-Function Studies of NSS-Proteins Integrates Information from Bioinformatics and Biomedical Literature 2010 ,		1
25	A Structural Context for Studying Neurotransmitter Transporter Function 2004 , 213-234		1
24	Distinct antagonist-bound inactive states underlie the divergence in the structures of the dopamine D2 and D3 receptors		1
23	Conformational Plasticity of GPCR Binding Sites 2005 , 363-388		1
22	A novel luminescence-based Earrestin membrane recruitment assay for unmodified GPCRs		1
21	Getting to grips with ammonium. <i>ELife</i> , 2013 , 2, e01029	8.9	1
20	Getting to grips with ammonium. <i>ELife</i> , 2013 , 2, e01029 Novel Class of Psychedelic Iboga Alkaloids Disrupts Opioid Addiction States	8.9	1
		8.9 4.6	
20	Novel Class of Psychedelic Iboga Alkaloids Disrupts Opioid Addiction States The Role of the Dopamine Transporter in the Effects of Amphetamine on Sleep and Sleep	4.6	1
20	Novel Class of Psychedelic Iboga Alkaloids Disrupts Opioid Addiction States The Role of the Dopamine Transporter in the Effects of Amphetamine on Sleep and Sleep Architecture in Drosophila. <i>Neurochemical Research</i> , 2021 , 1 Reply to Pantipsychotics with similar association kinetics at dopamine D receptors differ in	4.6	1
20 19 18	Novel Class of Psychedelic Iboga Alkaloids Disrupts Opioid Addiction States The Role of the Dopamine Transporter in the Effects of Amphetamine on Sleep and Sleep Architecture in Drosophila. <i>Neurochemical Research</i> , 2021 , 1 Reply to Pantipsychotics with similar association kinetics at dopamine D receptors differ in extrapyramidal side-effects. <i>Nature Communications</i> , 2018 , 9, 3568 Tianeptine, but not fluoxetine, decreases avoidant behavior in a mouse model of early	4.6	1 1
20 19 18	Novel Class of Psychedelic Iboga Alkaloids Disrupts Opioid Addiction States The Role of the Dopamine Transporter in the Effects of Amphetamine on Sleep and Sleep Architecture in Drosophila. <i>Neurochemical Research</i> , 2021 , 1 Reply to Pantipsychotics with similar association kinetics at dopamine D receptors differ in extrapyramidal side-effects P. <i>Nature Communications</i> , 2018 , 9, 3568 Tianeptine, but not fluoxetine, decreases avoidant behavior in a mouse model of early developmental exposure to fluoxetine. <i>Scientific Reports</i> , 2021 , 11, 22852	4.6 17.4 4.9	1 1 1
20 19 18 17 16	Novel Class of Psychedelic Iboga Alkaloids Disrupts Opioid Addiction States The Role of the Dopamine Transporter in the Effects of Amphetamine on Sleep and Sleep Architecture in Drosophila. <i>Neurochemical Research</i> , 2021 , 1 Reply to Pantipsychotics with similar association kinetics at dopamine D receptors differ in extrapyramidal side-effects. <i>Nature Communications</i> , 2018 , 9, 3568 Tianeptine, but not fluoxetine, decreases avoidant behavior in a mouse model of early developmental exposure to fluoxetine. <i>Scientific Reports</i> , 2021 , 11, 22852 Assays for detecting arrestin interaction with GPCRs. <i>Methods in Cell Biology</i> , 2021 , 166, 43-65 OZITX, a pertussis toxin-like protein for occluding inhibitory G protein signalling including GD	4.6 17.4 4.9 1.8	1 1 0 0

Substituted Cysteine Accessibility Method (SCAM)229-250

11	The Binding Pocket of G-Protein-Coupled Receptors for Biogenic Amines, Retinal, and Other Ligands 2003 , 155-160	
10	Transmembrane five effects on functional selectivity at the dopamine D2L receptor. <i>FASEB Journal</i> , 2006 , 20, A246	0.9
9	Identification of intracellular residues in the dopamine transporter critical for regulation of transporter conformation and cocaine binding. VOLUME 279 (2004) PAGES 3228-3238. <i>Journal of Biological Chemistry</i> , 2006 , 281, 25867-25868	5.4
8	Imaging Functional Dynamic Processes within Integral Membrane Proteins at the Single-Molecule Scale. <i>FASEB Journal</i> , 2015 , 29, 498.3	0.9
7	Lipid rafts and membrane cholesterol are involved in regulating D2 dopamine receptor signaling. <i>FASEB Journal</i> , 2010 , 24, 584.1	0.9
6	Towards Better Understanding of G(s) Coupling in Catecholamine Receptors 2014, 89-90	
5	The Membrane-Raft Protein Flotillin-1 is Essential in Dopamine Neurons for Amphetamine-Induced Behavior in Drosophila 2014 , 58	
4	Deciphering the Functionally Selective Properties of D2R Ligands 2014 , 110	
3	High-Throughput Screening for Modulators of the D2 Dopamine Receptor Yields Unique and Selective Pharmacological Chemotypes 2014 , 115	
2	Encephalopathy-causing mutations in GI() alter regulation of neuronal GIRK channels. <i>IScience</i> , 2021 , 24, 103018	6.1
1	Functional Genomic Analysis of Amphetamine Sensitivity in Frontiers in Psychiatry, 2022, 13, 831597	5