Jonathan A Javitch

List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/1146698/jonathan-a-javitch-publications-by-year.pdf

Version: 2024-04-28

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

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

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

#	Paper	IF	Citations
263	Functional Genomic Analysis of Amphetamine Sensitivity in Frontiers in Psychiatry, 2022, 13, 831597	5	
262	OZITX, a pertussis toxin-like protein for occluding inhibitory G protein signalling including GII <i>Communications Biology</i> , 2022 , 5, 256	6.7	0
261	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
260	Tianeptine, but not fluoxetine, decreases avoidant behavior in a mouse model of early developmental exposure to fluoxetine. <i>Scientific Reports</i> , 2021 , 11, 22852	4.9	0
259	Dopamine D2 receptors modulate the cholinergic pause and inhibitory learning. <i>Molecular Psychiatry</i> , 2021 ,	15.1	3
258	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
257	Disrupting D1-NMDA or D2-NMDA receptor heteromerization prevents cocaineß rewarding effects but preserves natural reward processing. <i>Science Advances</i> , 2021 , 7, eabg5970	14.3	3
256	Assays for detecting arrestin interaction with GPCRs. Methods in Cell Biology, 2021, 166, 43-65	1.8	0
255	Single-molecule FRET imaging of GPCR dimers in living cells. <i>Nature Methods</i> , 2021 , 18, 397-405	21.6	30
254	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
253	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
252	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
251	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
250	Crystal structures of LeuT reveal conformational dynamics in the outward-facing states. <i>Journal of Biological Chemistry</i> , 2021 , 296, 100609	5.4	3
249	Controlling opioid receptor functional selectivity by targeting distinct subpockets of the orthosteric site. <i>ELife</i> , 2021 , 10,	8.9	16
248	The Role of the Dopamine Transporter in the Effects of Amphetamine on Sleep and Sleep Architecture in Drosophila. <i>Neurochemical Research</i> , 2021 , 1	4.6	1
247	How changes in dopamine D2 receptor levels alter striatal circuit function and motivation. Molecular Psychiatry, 2021,	15.1	2

(2020-2021)

246	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
245	Encephalopathy-causing mutations in G[]) alter regulation of neuronal GIRK channels. <i>IScience</i> , 2021 , 24, 103018	6.1	
244	Mu opioid receptors on hippocampal GABAergic interneurons are critical for the antidepressant effects of tianeptine. <i>Neuropsychopharmacology</i> , 2021 ,	8.7	3
243	Cortical overgrowth in a preclinical forebrain organoid model of CNTNAP2-associated autism spectrum disorder. <i>Nature Communications</i> , 2021 , 12, 4087	17.4	5
242	A novel luminescence-based Enrestin recruitment assay for unmodified receptors. <i>Journal of Biological Chemistry</i> , 2021 , 296, 100503	5.4	5
241	Structure of human GABA receptor in an inactive state. <i>Nature</i> , 2020 , 584, 304-309	50.4	32
240	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
239	Signalling profiles of a structurally diverse panel of synthetic cannabinoid receptor agonists. <i>Biochemical Pharmacology</i> , 2020 , 175, 113871	6	15
238	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
237	Ribosome-associated vesicles: A dynamic subcompartment of the endoplasmic reticulum in secretory cells. <i>Science Advances</i> , 2020 , 6, eaay9572	14.3	20
236	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
235	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
234	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
233	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
232	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
231	Novel Fluorescent Ligands Enable Single-Molecule Localization Microscopy of the Dopamine Transporter. <i>ACS Chemical Neuroscience</i> , 2020 , 11, 3288-3300	5.7	3
230	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
229	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

228	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
227	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
226	Arrestin recruitment to dopamine D2 receptor mediates locomotion but not incentive motivation. <i>Molecular Psychiatry</i> , 2020 , 25, 2086-2100	15.1	32
225	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
224	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
223	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
222	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
221	Cannabinoid CB1 and CB2 Receptor-Mediated Arrestin Translocation: Species, Subtype, and Agonist-Dependence. <i>Frontiers in Pharmacology</i> , 2019 , 10, 350	5.6	26
220	Molecular Determinants of the Intrinsic Efficacy of the Antipsychotic Aripiprazole. <i>ACS Chemical Biology</i> , 2019 , 14, 1780-1792	4.9	14
219	Quantifying secondary transport at single-molecule resolution. <i>Nature</i> , 2019 , 575, 528-534	50.4	22
218	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, 159	47-1:59	56 ⁶
217	Role of Tau Protein in Remodeling of Circadian Neuronal Circuits and Sleep. <i>Frontiers in Aging Neuroscience</i> , 2019 , 11, 320	5.3	11
216	Regional Heterogeneity of D2-Receptor Signaling in the Dorsal Striatum and Nucleus Accumbens. <i>Neuron</i> , 2018 , 98, 575-587.e4	13.9	41
215	Gs-Iversus Golf-dependent functional selectivity mediated by the dopamine D receptor. <i>Nature Communications</i> , 2018 , 9, 486	17.4	24
214	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
213	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
212	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
211	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

(2017-2018)

210	Accumbens dopamine D2 receptors increase motivation by decreasing inhibitory transmission to the ventral pallidum. <i>Nature Communications</i> , 2018 , 9, 1086	17.4	48
209	Treatment resistant depression: A multi-scale, systems biology approach. <i>Neuroscience and Biobehavioral Reviews</i> , 2018 , 84, 272-288	9	209
208	Luciferase complementation based-detection of G-protein-coupled receptor activity. <i>BioTechniques</i> , 2018 , 65, 9-14	2.5	10
207	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
206	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
205	Reply to PAntipsychotics with similar association kinetics at dopamine D receptors differ in extrapyramidal side-effects. <i>Nature Communications</i> , 2018 , 9, 3568	17.4	1
204	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
203	Potentiating SLC transporter activity: Emerging drug discovery opportunities. <i>Biochemical Pharmacology</i> , 2017 , 135, 1-11	6	35
202	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	-456	36
201	Phospho-specific antibodies targeting the amino terminus of the human dopamine transporter. Journal of Chemical Neuroanatomy, 2017 , 83-84, 91-98	3.2	4
200	Single-molecule analysis of ligand efficacy in AR-G-protein activation. <i>Nature</i> , 2017 , 547, 68-73	50.4	164
199	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
198	The Behavioral Effects of the Antidepressant Tianeptine Require the Mu-Opioid Receptor. Neuropsychopharmacology, 2017 , 42, 2052-2063	8.7	123
197	Toward Understanding the Structural Basis of Partial Agonism at the Dopamine D Receptor. Journal of Medicinal Chemistry, 2017 , 60, 580-593	8.3	33
196	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
195	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
194	Extreme Vetting of Dopamine Receptor Oligomerization 2017, 99-127		2
193	Neuronal Depolarization Drives Increased Dopamine Synaptic Vesicle Loading via VGLUT. <i>Neuron</i> , 2017 , 95, 1074-1088.e7	13.9	42

192	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
191	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
190	Electronic tuning of self-healing fluorophores for live-cell and single-molecule imaging. <i>Chemical Science</i> , 2017 , 8, 755-762	9.4	36
189	Dopamine D2 Receptors in the Paraventricular Thalamus Attenuate Cocaine Locomotor Sensitization. <i>ENeuro</i> , 2017 , 4,	3.9	22
188	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
187	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
186	Development and Antiparkinsonian Activity of VU0418506, a Selective Positive Allosteric Modulator of Metabotropic Glutamate Receptor 4 Homomers without Activity at mGlu2/4 Heteromers. ACS Chemical Neuroscience, 2016, 7, 1201-11	5.7	47
185	The role of kinetic context in apparent biased agonism at GPCRs. <i>Nature Communications</i> , 2016 , 7, 1084	1217.4	206
184	Role of Annular Lipids in the Functional Properties of Leucine Transporter LeuT Proteomicelles. <i>Biochemistry</i> , 2016 , 55, 850-9	3.2	10
183	Development of a Rapid Insulin Assay by Homogenous Time-Resolved Fluorescence. <i>PLoS ONE</i> , 2016 , 11, e0148684	3.7	18
183		3.7 8.3	18
	2016, 11, e0148684 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</i>		26
182	2016, 11, e0148684 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 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</i>	8.3	26
182	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 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 Dual agonist occupancy of AT1-R-DC-AR heterodimers results in atypical Gs-PKA signaling. <i>Nature</i>	8.3	26 161
182 181 180	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 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 Dual agonist occupancy of AT1-R-DC-AR heterodimers results in atypical Gs-PKA signaling. <i>Nature Chemical Biology</i> , 2015 , 11, 271-9 Evidence for limited D1 and D2 receptor coexpression and colocalization within the dorsal striatum	8.3	26 161 76
182 181 180	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 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 Dual agonist occupancy of AT1-R-2C-AR heterodimers results in atypical Gs-PKA signaling. <i>Nature Chemical Biology</i> , 2015, 11, 271-9 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 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,	8.3 16.4 11.7	261617621
182 181 180 179	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 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 Dual agonist occupancy of AT1-R-2C-AR heterodimers results in atypical Gs-PKA signaling. <i>Nature Chemical Biology</i> , 2015 , 11, 271-9 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 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 Upregulation of dopamine D2 receptors in the nucleus accumbens indirect pathway increases	8.3 16.4 11.7 3.4 8.3	26161762135

(2014-2015)

174	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	
173	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	
172	Evidence against dopamine D1/D2 receptor heteromers. <i>Molecular Psychiatry</i> , 2015 , 20, 1373-85	15.1	83	
171	Imaging Functional Dynamic Processes within Integral Membrane Proteins at the Single-Molecule Scale. <i>FASEB Journal</i> , 2015 , 29, 498.3	0.9		
170	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	
169	Identification of novel functionally selective Eppioid receptor scaffolds. <i>Molecular Pharmacology</i> , 2014 , 85, 83-90	4.3	95	
168	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	
167	A new mechanism of allostery in a G protein-coupled receptor dimer. <i>Nature Chemical Biology</i> , 2014 , 10, 745-52	11.7	95	
166	The atypical antidepressant and neurorestorative agent tianeptine is a Eppioid receptor agonist. <i>Translational Psychiatry</i> , 2014 , 4, e411	8.6	74	
165	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	
164	PIP2 regulates psychostimulant behaviors through its interaction with a membrane protein. <i>Nature Chemical Biology</i> , 2014 , 10, 582-589	11.7	83	
163	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	
162	Conformational dynamics of ligand-dependent alternating access in LeuT. <i>Nature Structural and Molecular Biology</i> , 2014 , 21, 472-9	17.6	102	
161	Cross-Talk between G Protein-Coupled Receptors: Challenges of Distinguishing Upstream from Downstream Mechanisms 2014 , 93-94			
160	Rebuttal from Nevin A. Lambert and Jonathan A. Javitch. <i>Journal of Physiology</i> , 2014 , 592, 2449	3.9	12	
159	Conformational changes in dopamine transporter intracellular regions upon cocaine binding and dopamine translocation. <i>Neurochemistry International</i> , 2014 , 73, 4-15	4.4	10	
158	Dopamine receptor activation increases HIV entry into primary human macrophages. <i>PLoS ONE</i> , 2014 , 9, e108232	3.7	45	
157	Towards Better Understanding of G(s) Coupling in Catecholamine Receptors 2014 , 89-90			

The Membrane-Raft Protein Flotillin-1 is Essential in Dopamine Neurons for Amphetamine-Induced Behavior in Drosophila **2014**, 58

155	Deciphering the Functionally Selective Properties of D2R Ligands 2014 , 110		
154	High-Throughput Screening for Modulators of the D2 Dopamine Receptor Yields Unique and Selective Pharmacological Chemotypes 2014 , 115		
153	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
152	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
151	Increasing dopamine D2 receptor expression in the adult nucleus accumbens enhances motivation. <i>Molecular Psychiatry</i> , 2013 , 18, 1025-33	15.1	137
150	Segregation of family A G protein-coupled receptor protomers in the plasma membrane. <i>Molecular Pharmacology</i> , 2013 , 84, 346-52	4.3	25
149	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
148	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
147	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
146	Getting to grips with ammonium. <i>ELife</i> , 2013 , 2, e01029	8.9	1
145	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
144	Dopamine-mediated autocrine inhibitory circuit regulating human insulin secretion in vitro. <i>Molecular Endocrinology</i> , 2012 , 26, 1757-72		61
143	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
142	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
141	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
140	Molecular determinants of selectivity and efficacy at the dopamine D3 receptor. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 6689-99	8.3	131
139	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

138	Cyanine fluorophore derivatives with enhanced photostability. <i>Nature Methods</i> , 2011 , 9, 68-71	21.6	203
137	CODA-RET reveals functional selectivity as a result of GPCR heteromerization. <i>Nature Chemical Biology</i> , 2011 , 7, 624-30	11.7	92
136	Making structural sense of dimerization interfaces of delta opioid receptor homodimers. <i>Biochemistry</i> , 2011 , 50, 1682-90	3.2	66
135	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
134	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
133	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
132	Crystal structure of a potassium ion transporter, TrkH. <i>Nature</i> , 2011 , 471, 336-40	50.4	101
131	Crystal structure of a phosphorylation-coupled saccharide transporter. <i>Nature</i> , 2011 , 473, 50-4	50.4	67
130	Substrate-modulated gating dynamics in a Na+-coupled neurotransmitter transporter homologue. <i>Nature</i> , 2011 , 474, 109-13	50.4	244
129	Chapter 12:Crosstalk Between Receptors: Challenges of Distinguishing Upstream from Downstream Mechanisms. <i>RSC Drug Discovery Series</i> , 2011 , 255-268	0.6	
128	Discovery of Enrestin-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
127	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
126	Presynaptic regulation of dopamine transmission in schizophrenia. Schizophrenia Bulletin, 2011, 37, 108	-173	47
125	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
124	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
123	Single-molecule dynamics of gating in a neurotransmitter transporter homologue. <i>Nature</i> , 2010 , 465, 188-93	50.4	213
122	Substrate-dependent proton antiport in neurotransmitter:sodium symporters. <i>Nature Chemical Biology</i> , 2010 , 6, 109-16	11.7	48
121	Time-resolved FRET between GPCR ligands reveals oligomers in native tissues. <i>Nature Chemical Biology</i> , 2010 , 6, 587-94	11.7	277

120	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
119	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
118	GPCR-OKB: the G Protein Coupled Receptor Oligomer Knowledge Base. <i>Bioinformatics</i> , 2010 , 26, 1804-5	57.2	71
117	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
116	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
115	Akt-dependent and isoform-specific regulation of dopamine transporter cell surface expression. <i>ACS Chemical Neuroscience</i> , 2010 , 1, 476-81	5.7	24
114	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
113	Signaling pathways in schizophrenia: emerging targets and therapeutic strategies. <i>Trends in Pharmacological Sciences</i> , 2010 , 31, 381-90	13.2	128
112	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
111	Impact of D2 receptor internalization on binding affinity of neuroimaging radiotracers. <i>Neuropsychopharmacology</i> , 2010 , 35, 806-17	8.7	63
110	TRAC: A Platform for Structure-Function Studies of NSS-Proteins Integrates Information from Bioinformatics and Biomedical Literature 2010 ,		1
109	Structural Basis of Dopamine Receptor Activation 2010 , 47-73		3
108	Lipid rafts and membrane cholesterol are involved in regulating D2 dopamine receptor signaling. <i>FASEB Journal</i> , 2010 , 24, 584.1	0.9	
107	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
106	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
105	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
104	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-	.155 ⁴	58
103	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

(2007-2009)

102	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
101	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
100	Allosteric communication between protomers of dopamine class A GPCR dimers modulates activation. <i>Nature Chemical Biology</i> , 2009 , 5, 688-95	11.7	294
99	Building a new conceptual framework for receptor heteromers. <i>Nature Chemical Biology</i> , 2009 , 5, 131-4	11.7	313
98	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
97	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
96	Drs. Lieberman, Javitch, and Moore Reply. American Journal of Psychiatry, 2009, 166, 111-113	11.9	3
95	Dopamine D2 receptors form higher order oligomers at physiological expression levels. <i>EMBO Journal</i> , 2008 , 27, 2293-304	13	286
94	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
93	The binding sites for cocaine and dopamine in the dopamine transporter overlap. <i>Nature Neuroscience</i> , 2008 , 11, 780-9	25.5	260
92	Syntaxin 1A interaction with the dopamine transporter promotes amphetamine-induced dopamine efflux. <i>Molecular Pharmacology</i> , 2008 , 74, 1101-8	4.3	98
91	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
90	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
89	PI3K signaling supports amphetamine-induced dopamine efflux. <i>Biochemical and Biophysical Research Communications</i> , 2008 , 372, 656-61	3.4	34
88	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
87	An intracellular interaction network regulates conformational transitions in the dopamine transporter. <i>Journal of Biological Chemistry</i> , 2008 , 283, 17691-701	5.4	108
86	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
85	Mechanism of chloride interaction with neurotransmitter:sodium symporters. <i>Nature</i> , 2007 , 449, 726-30) 50.4	188

84	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
83	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
82	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
81	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
80	Functional selectivity and classical concepts of quantitative pharmacology. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2007 , 320, 1-13	4.7	870
79	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
78	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
77	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. <i>Molecular Pharmacology</i> , 2006 , 70, 1630-42	4.3	232
76	Regulation of dopamine transporter trafficking by intracellular amphetamine. <i>Molecular Pharmacology</i> , 2006 , 70, 542-8	4.3	57
75	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
75 74		0.9	177
	amphetamine-induced reverse transport. <i>Neuron</i> , 2006 , 51, 417-29 Transmembrane five effects on functional selectivity at the dopamine D2L receptor. <i>FASEB Journal</i> ,		177
74	amphetamine-induced reverse transport. <i>Neuron</i> , 2006 , 51, 417-29 Transmembrane five effects on functional selectivity at the dopamine D2L receptor. <i>FASEB Journal</i> , 2006 , 20, A246 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</i>	0.9	217
74 73	amphetamine-induced reverse transport. <i>Neuron</i> , 2006 , 51, 417-29 Transmembrane five effects on functional selectivity at the dopamine D2L receptor. <i>FASEB Journal</i> , 2006 , 20, A246 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 Amphetamine induces dopamine efflux through a dopamine transporter channel. <i>Proceedings of</i>	o.9 5·4	
74 73 72	Transmembrane five effects on functional selectivity at the dopamine D2L receptor. <i>FASEB Journal</i> , 2006 , 20, A246 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 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	o.9 5·4	217
74 73 72 71	Transmembrane five effects on functional selectivity at the dopamine D2L receptor. <i>FASEB Journal</i> , 2006 , 20, A246 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 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 Oligomerization Domains of G Protein-Coupled Receptors 2005 , 243-265 Intramolecular cross-linking in a bacterial homolog of mammalian SLC6 neurotransmitter transporters suggests an evolutionary conserved role of transmembrane segments 7 and 8.	0.9 5.4 11.5	217
74 73 72 71 70	Transmembrane five effects on functional selectivity at the dopamine D2L receptor. <i>FASEB Journal</i> , 2006 , 20, A246 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 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 Oligomerization Domains of G Protein-Coupled Receptors 2005 , 243-265 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 A pincer-like configuration of TM2 in the human dopamine transporter is responsible for indirect	0.9 5.4 11.5	217 4 11

(2003-2005)

66	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
65	Conformational Plasticity of GPCR Binding Sites 2005 , 363-388		1
64	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
63	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
62	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
61	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
60	The second extracellular loop of the dopamine D2 receptor lines the binding-site crevice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2004 , 101, 440-5	11.5	198
59	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
58	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
57	N-terminal phosphorylation of the dopamine transporter is required for amphetamine-induced efflux. <i>PLoS Biology</i> , 2004 , 2, E78	9.7	188
56	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
55	A Structural Context for Studying Neurotransmitter Transporter Function 2004 , 213-234		1
54	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
53	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
52	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
51	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
50	Probing conformational changes in neurotransmitter transporters: a structural context. <i>European Journal of Pharmacology</i> , 2003 , 479, 3-12	5.3	36
49	The fourth transmembrane segment forms the interface of the dopamine D2 receptor homodimer. Journal of Biological Chemistry, 2003 , 278, 4385-8	5.4	242

48	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
47	Characterization of a functional bacterial homologue of sodium-dependent neurotransmitter transporters. <i>Journal of Biological Chemistry</i> , 2003 , 278, 12703-9	5.4	77
46	The Binding Pocket of G-Protein-Coupled Receptors for Biogenic Amines, Retinal, and Other Ligands 2003 , 155-160		
45	PI 3-kinase regulation of dopamine uptake. <i>Journal of Neurochemistry</i> , 2002 , 81, 859-69	6	168
44	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
43	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
42	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
41	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
40	Zn(2+) site engineering at the oligomeric interface of the dopamine transporter. <i>FEBS Letters</i> , 2002 , 524, 87-91	3.8	31
39	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
38	Reaction of oxidized dopamine with endogenous cysteine residues in the human dopamine transporter. <i>Journal of Neurochemistry</i> , 2001 , 76, 1242-51	6	158
37	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
36	[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
35	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
34	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
33	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
32	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
31	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

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
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
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
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
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
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
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
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
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
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
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
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
Mapping the binding-site crevice of the D2 receptor. Advances in Pharmacology, 1998, 42, 412-5	5.7	13
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
Probing structure of neurotransmitter transporters by substituted-cysteine accessibility method. <i>Methods in Enzymology</i> , 1998 , 296, 331-46	1.7	46
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
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
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
	The substituted-cysteine accessibility method (SCAM) to elucidate membrane protein structure. Current Protocols in Neuroscience, 2001, Chapter 4, Unit 4.15 Structural mimicry in G protein-coupled receptors: implications of the high-resolution structure of rhodopsin for structure-function analysis of rhodopsin-like receptors. Molecular Pharmacology, 2001, 60, 1-19 The forgotten serine. A critical role for Ser-2035 42 in ligand binding to and activation of the beta 2-adrenergic receptor. Journal of Biological Chemistry, 2000, 275, 37779-88 Transport-dependent accessibility of a cytoplasmic loop cysteine in the human dopamine transporter. Journal of Biological Chemistry, 2000, 275, 1608-14 Amphetamine-induced loss of human dopamine transporter activity: an internalization-dependent and cocaine-sensitive mechanism. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 6850-5 The conserved cysteine 7.38 residue is differentially accessible in the binding-site crevices of the mu, delta, and kappa opioid receptors. Biochemistry, 2000, 39, 13904-15 The fourth transmembrane segment of the dopamine D2 receptor: accessibility in the binding-site crevice and position in the transmembrane bundle. Biochemistry, 2000, 39, 12190-9 Dopamine D4/D2 receptor selectivity is determined by A divergent aromatic microdomain contained within the second, third, and seventh membrane-spanning segments. Molecular Pharmacology, 1999, 56, 1116-26 Mutation of a highly conserved aspartic acid in the beta2 adrenergic receptor: constitutive activation, structural instability, and conformational rearrangement of transmembrane segment 6. Molecular Pharmacology, 1999, 56, 1178-84 Chloroethylclonidine and 2-aminoethyl methanethiosulfonate recognize two different conformations of the human alpha(2A)-adrenergic receptor. Journal of Biological Chemistry, 1999, 274, 21867-72 Electrostatic and aromatic microdomains within the binding-site crevice of the D2 receptor: contributions of the second membrane-spann	The substituted-cysteine accessibility method (SCAM) to elucidate membrane protein structure. Current Protocols in Neuroscience, 2001, Chapter 4, Unit 4.15 Structural mimicry in G protein-coupled receptors: implications of the high-resolution structure of rhodopsin for structure-function analysis of rhodopsin-like receptors. Molecular Pharmacology, 2001, 60, 1-19 The forgotten serine. A critical role for Ser-2035.42 in ligand binding to and activation of the beta 2-adrenergic receptor. Journal of Biological Chemistry, 2000, 275, 37779-88 Transport-dependent accessibility of a cytoplasmic loop cysteine in the human dopamine transporter. Journal of Biological Chemistry, 2000, 275, 1608-14 Amphetamine-induced loss of human dopamine transporter activity: an internalization-dependent and cocaine-sensitive mechanism. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 6850-5 The conserved cysteine 7.38 residue is differentially accessible in the binding-site crevices of the mu, delta, and kappa opioid receptors. Biochemistry, 2000, 39, 13904-15 The fourth transmembrane segment of the dopamine D2 receptor: accessibility in the binding-site crevice and position in the transmembrane bundle. Biochemistry, 2000, 39, 12190-9 Dopamine D4/D2 receptor selectivity is determined by A divergent aromatic microdomain contained within the second, third, and seventh membrane-spanning segments. Molecular Pharmacology, 1999, 56, 1716-26 Mutation of a highly conserved asparatic acid in the beta2 adrenergic receptor: constitutive activation, structural instability, and conformational rearrangement of transmembrane segment 6. Molecular Pharmacology, 1999, 56, 175-84 Chloroethylclonidine and 2-aminoethyl methanethiosulfonate recognize two different conformations of the human alpha(2A)-adrenergic receptor. Journal of Biological Chemistry, 1999, 38, 7961-8 Mapping the binding-site crevice of the D2 receptor. Advances in Pharmacology, 1998, 42, 412-5 A cluster of aromatic residues in the si

12	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
11	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
10	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
9	Differential visualization of dopamine D2 and D3 receptors in rat brain. <i>European Journal of Pharmacology</i> , 1993 , 234, 269-72	5.3	15
8	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
7	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
6	[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
5	Substituted Cysteine Accessibility Method (SCAM)229-250		
4	Distinct antagonist-bound inactive states underlie the divergence in the structures of the dopamine D2 and D3 receptors		1
3	A novel luminescence-based Earrestin membrane recruitment assay for unmodified GPCRs		1
2	Dopamine D2 receptors modulate the cholinergic pause and inhibitory learning		2
1	Novel Class of Psychedelic Iboga Alkaloids Disrupts Opioid Addiction States		1