Jonathan B Cohen

List of Publications by Year
in descending order

98	5,306	66234	85405
papers	citations	42 h-index	71 g-index

101	101	101	1959
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Photoaffinity labeling identifies an intersubunit steroid-binding site in heteromeric GABA type A (GABAA) receptors. Journal of Biological Chemistry, 2020, 295, 11495-11512.	1.6	10
2	A potent photoreactive general anesthetic with novel binding site selectivity for GABAA receptors. European Journal of Medicinal Chemistry, 2020, 194, 112261.	2.6	3
3	Competitive Antagonism of Etomidate Action by Diazepam. Anesthesiology, 2020, 133, 583-594.	1.3	7
4	Identifying Drugs that Bind Selectively to Intersubunit General Anesthetic Sites in the Î±1Î²3Î³2 GABA_AR Transmembrane Domain. Molecular Pharmacology, 2019, 95, 615-628.	1.0	22
5	A photoreactive analog of allopregnanolone enables identification of steroid-binding sites in a nicotinic acetylcholine receptor. Journal of Biological Chemistry, 2019, 294, 7892-7903.	1.6	3
6	Inhibitable photolabeling by neurosteroid diazirine analog in the Î²3-Subunit of human hetereopentameric type A GABA receptors. European Journal of Medicinal Chemistry, 2019, 162, 810-824.	2.6	7
7	Etomidate and Etomidate Analog Binding and Positive Modulation of Î³-Aminobutyric Acid Type A Receptors. Anesthesiology, 2018, 129, 959-969.	1.3	8
8	Enantiomeric barbiturates bind distinct inter- and intrasubunit binding sites in a nicotinic acetylcholine receptor (nAChR). Journal of Biological Chemistry, 2017, 292, 17258-17271.	1.6	1
9	Competitive Antagonism of Anesthetic Action at the Î³-Aminobutyric Acid Type A Receptor by a Novel Etomidate Analog with Low Intrinsic Efficacy. Anesthesiology, 2017, 127, 824-837.	1.3	9
10	Synthesis and pharmacological evaluation of neurosteroid photoaffinity ligands. European Journal of Medicinal Chemistry, 2017, 136, 334-347.	2.6	12
11	Photolabeling a Nicotinic Acetylcholine Receptor (nAChR) with an (Î±4)₃(Î²2)₂ nAChR-Selective Positive Allosteric Modulator. Molecular Pharmacology, 2016, 89, 575-584.	1.0	15
12	Interactions of a Photoreactive Steroid Anesthetic (F4N3-Alphaxalone) with Human Î±1Î²3Î³2 GABA-A Receptors. Biophysical Journal, 2016, 110, 455a.	0.2	1
13	General Anesthetic Binding Sites in Human Î±4Î²3Î³ Î³-Aminobutyric Acid Type A Receptors (GABAARs). Journal of Biological Chemistry, 2016, 291, 26529-26539.	1.6	19
14	Multiple Non-Equivalent Interfaces Mediate Direct Activation of GABAA Receptors by Propofol. Current Neuropharmacology, 2016, 14, 772-780.	1.4	37
15	Desformylflustrabromine (dFBr) and [³H]dFBr-Labeled Binding Sites in a Nicotinic Acetylcholine Receptor. Molecular Pharmacology, 2015, 88, 1-11.	1.0	22
16	Positive and Negative Allosteric Modulation of an Î±1Î²3Î³2 Î³-Aminobutyric Acid Type A (GABAA) Receptor by Binding to a Site in the Transmembrane Domain at the Î³+-Î²â^' Interface. Journal of Biological Chemistry, 2015, 290, 23432-23446.	1.6	28
17	Identifying Barbiturate Binding Sites in a Nicotinic Acetylcholine Receptor with [³H]Allyl m-Trifluoromethyldiazirine Mephobarbital, a Photoreactive Barbiturate. Molecular Pharmacology, 2014, 85, 735-746.	1.0	23
18	Photoaffinity Labeling of Nicotinic Receptors: Diversity of Drug Binding Sites!. Journal of Molecular Neuroscience, 2014, 53, 480-486.	1.1	32

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19	Multiple Propofol-binding Sites in a Î³-Aminobutyric Acid Type A Receptor (GABAAR) Identified Using a Photoreactive Propofol Analog. Journal of Biological Chemistry, 2014, 289, 27456-27468.	1.6	106
20	Photoaffinity Labeling the Propofol Binding Site in GLIC. Biochemistry, 2014, 53, 135-142.	1.2	36
21	Specificity of Intersubunit General Anesthetic-binding Sites in the Transmembrane Domain of the Human Î±1Î²3Î³2 Î³-Aminobutyric Acid Type A (GABAA) Receptor*. Journal of Biological Chemistry, 2013, 288, 19343-19357.	1.6	124
22	Identification of Propofol Binding Sites in a Nicotinic Acetylcholine Receptor with a Photoreactive Propofol Analog*. Journal of Biological Chemistry, 2013, 288, 6178-6189.	1.6	69
23	Physostigmine and Galanthamine Bind in the Presence of Agonist at the Canonical and Noncanonical Subunit Interfaces of a Nicotinic Acetylcholine Receptor. Journal of Neuroscience, 2013, 33, 485-494.	1.7	49
24	Bupropion Binds to Two Sites in theTorpedoNicotinic Acetylcholine Receptor Transmembrane Domain: A Photoaffinity Labeling Study with the Bupropion Analogue [125I]-SADU-3-72. Biochemistry, 2012, 51, 2425-2435.	1.2	24
25	Mapping General Anesthetic Binding Site(s) in Human Î±1Î²3 Î³-Aminobutyric Acid Type A Receptors with [³H]TDBzl-Etomidate, a Photoreactive Etomidate Analogue. Biochemistry, 2012, 51, 836-847.	1.2	98
26	Allylm-Trifluoromethyldiazirine Mephobarbital: An Unusually Potent Enantioselective and Photoreactive Barbiturate General Anesthetic. Journal of Medicinal Chemistry, 2012, 55, 6554-6565.	2.9	46
27	Identifying an Etomidate Binding Site in Heterologously Expressed Human Alpha1/Beta3 GABAA Receptors (GABAAR) Using Photoactive Etomidate Analogs. Biophysical Journal, 2011, 100, 271a.	0.2	1
28	p-(4-Azipentyl)propofol: A Potent Photoreactive General Anesthetic Derivative of Propofol. Journal of Medicinal Chemistry, 2011, 54, 8124-8135.	2.9	35
29	Multiple Transmembrane Binding Sites for p-Trifluoromethyldiazirinyl-etomidate, a Photoreactive Torpedo Nicotinic Acetylcholine Receptor Allosteric Inhibitor. Journal of Biological Chemistry, 2011, 286, 20466-20477.	1.6	36
30	Conformational Changes in the Nicotinic Acetylcholine Receptor during Gating and Desensitization. Biochemistry, 2010, 49, 156-165.	1.2	34
31	p-Trifluoromethyldiazirinyl-etomidate: A Potent Photoreactive General Anesthetic Derivative of Etomidate That Is Selective for Ligand-Gated Cationic Ion Channels. Journal of Medicinal Chemistry, 2010, 53, 6432-6444.	2.9	24
32	Numerous Classes of General Anesthetics Inhibit Etomidate Binding to Î³-Aminobutyric Acid Type A (GABAA) Receptors. Journal of Biological Chemistry, 2010, 285, 8615-8620.	1.6	75
33	[3H]Epibatidine Photolabels Non-equivalent Amino Acids in the Agonist Binding Site of Torpedo and Î±4Î²2 Nicotinic Acetylcholine Receptors. Journal of Biological Chemistry, 2009, 284, 24939-24947.	1.6	12
34	Time-Resolved Photolabeling of the Nicotinic Acetylcholine Receptor by [³H]Azietomidate, an Open-State Inhibitor. Molecular Pharmacology, 2009, 75, 1084-1095.	1.0	29
35	Hydrophobic Photolabeling Studies Identify the Lipidâ^'Protein Interface of the 5-HT_{3A} Receptor. Biochemistry, 2009, 48, 9278-9286.	1.2	11
36	[³H]Chlorpromazine Photolabeling of the Torpedo Nicotinic Acetylcholine Receptor Identifies Two State-Dependent Binding Sites in the Ion Channel. Biochemistry, 2009, 48, 10066-10077.	1.2	34

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37	Neurosteroids Allosterically Modulate Binding of the Anesthetic Etomidate to Î³-Aminobutyric Acid Type A Receptors. Journal of Biological Chemistry, 2009, 284, 11771-11775.	1.6	64
38	Probing the Structure of the Affinity-Purified and Lipid-Reconstituted Torpedo Nicotinic Acetylcholine Receptor. Biochemistry, 2008, 47, 12787-12794.	1.2	33
39	Identification of Binding Sites in the Nicotinic Acetylcholine Receptor for TDBzl-etomidate, a Photoreactive Positive Allosteric Effector. Journal of Biological Chemistry, 2008, 283, 22051-22062.	1.6	63
40	[³H]Benzophenone Photolabeling Identifies State-Dependent Changes in Nicotinic Acetylcholine Receptor Structure. Biochemistry, 2007, 46, 10296-10307.	1.2	30
41	Identifying the Lipidâ^'Protein Interface of the Î±4Î²2 Neuronal Nicotinic Acetylcholine Receptor:â€‰Hydrophobic Photolabeling Studies with 3-(Trifluoromethyl)-3-(m-[¹²⁵I]iodophenyl)diazirine. Biochemistry, 2007, 46, 13837-13846.	1.2	17
42	Synthesis of Trifluoromethylaryl Diazirine and Benzophenone Derivatives of Etomidate that Are Potent General Anesthetics and Effective Photolabels for Probing Sites on Ligand-Gated Ion Channels. Journal of Medicinal Chemistry, 2006, 49, 4818-4825.	2.9	43
43	Identification of a GABAA Receptor Anesthetic Binding Site at Subunit Interfaces by Photolabeling with an Etomidate Analog. Journal of Neuroscience, 2006, 26, 11599-11605.	1.7	280
44	Site Specificity of Agonist-Induced Opening and Desensitization of theTorpedocalifornicaNicotinic Acetylcholine Receptorâ€. Biochemistry, 2006, 45, 195-204.	1.2	18
45	Mapping the Structural Requirements for Nicotinic Acetylcholine Receptor Activation by Using Tethered Alkyltrimethylammonium Agonists and Antagonists. Biochemistry, 2006, 45, 10641-10653.	1.2	13
46	Cholesterol Interacts with Transmembrane Î±-Helices M1, M3, and M4 of theTorpedoNicotinic Acetylcholine Receptor:Â Photolabeling Studies Using [3H]Azicholesterolâ€. Biochemistry, 2006, 45, 976-986.	1.2	79
47	alpha-Conotoxin GI benzoylphenylalanine derivatives. 1H-NMR structures and photoaffinity labeling of the Torpedo californica nicotinic acetylcholine receptor. FEBS Journal, 2006, 273, 1373-1388.	2.2	17
48	Gating-enhanced Accessibility of Hydrophobic Sites within the Transmembrane Region of the Nicotinic Acetylcholine Receptor's Î'-Subunit. Journal of Biological Chemistry, 2005, 280, 13631-13640.	1.6	47
49	Cell-Surface MuSK Self-Association:Â a Crucial Role for the Putative Signal Sequenceâ€. Biochemistry, 2005, 44, 16229-16238.	1.2	2
50	Photolabeling the Torpedo Nicotinic Acetylcholine Receptor with 4-Azido-2,3,5,6-tetrafluorobenzoylcholine, a Partial Agonist. Biochemistry, 2005, 44, 13447-13456.	1.2	20
51	Identification of Binding Sites in the Nicotinic Acetylcholine Receptor for [3H]Azietomidate, a Photoactivatable General Anesthetic. Journal of Biological Chemistry, 2004, 279, 17640-17649.	1.6	71
52	Identification of Amino Acids in the Nicotinic Acetylcholine Receptor Agonist Binding Site and Ion Channel Photolabeled by 4-[(3-Trifluoromethyl)-3H-Diazirin-3-yl]Benzoylcholine, a Novel Photoaffinity Antagonistâ€. Biochemistry, 2003, 42, 271-283.	1.2	20
53	Identification of Nicotinic Acetylcholine Receptor Amino Acids Photolabeled by the Volatile Anesthetic Halothane. Biochemistry, 2003, 42, 13457-13467.	1.2	95
54	2-(3-Methyl-3H-diaziren-3-yl)ethyl 1-(1-phenylethyl)-1H-imidazole-5-carboxylate:Â A Derivative of the Stereoselective General Anesthetic Etomidate for Photolabeling Ligand-Gated Ion Channels. Journal of Medicinal Chemistry, 2003, 46, 1257-1265.	2.9	83

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55	Identification of the Bovine Î³-Aminobutyric Acid Type A Receptor Î± Subunit Residues Photolabeled by the Imidazobenzodiazepine [3H]Ro15-4513. Journal of Biological Chemistry, 2002, 277, 50036-50045.	1.6	49
56	Mapping the Agonist Binding Site of the Nicotinic Acetylcholine Receptor by Cysteine Scanning Mutagenesis: Antagonist Footprint and Secondary Structure Prediction. Molecular Pharmacology, 2002, 61, 463-472.	1.0	25
57	Identification and characterization of membrane-associated polypeptides in Torpedo nicotinic acetylcholine receptor-rich membranes by hydrophobic photolabeling. Biochimica Et Biophysica Acta - Biomembranes, 2001, 1512, 215-224.	1.4	14
58	Site of Resting State Inhibition of the Nicotinic Acetylcholine Receptor by a Hydrophobic Inhibitor. Biochemistry, 2001, 40, 296-304.	1.2	15
59	Interactions between 3-(Trifluoromethyl)-3-(m-[125I]iodophenyl)diazirine and Tetracaine, Phencyclidine, or Histrionicotoxin in theTorpedo Species Nicotinic Acetylcholine Receptor Ion Channel. Molecular Pharmacology, 2001, 59, 1514-1522.	1.0	13
60	Contributions of Torpedo Nicotinic Acetylcholine Receptor Î³Trp-55 and Î´Trp-57 to Agonist and Competitive Antagonist Function. Journal of Biological Chemistry, 2001, 276, 2417-2426.	1.6	40
61	Role of Rapsyn Tetratricopeptide Repeat and Coiled-coil Domains in Self-association and Nicotinic Acetylcholine Receptor Clustering. Journal of Biological Chemistry, 2001, 276, 7475-7483.	1.6	110
62	Interactions of the Rapsyn RING-H2 Domain with Dystroglycan. Journal of Biological Chemistry, 2001, 276, 24911-24917.	1.6	89
63	The Agrin/MuSK Signaling Pathway Is Spatially Segregated from the Neuregulin/ErbB Receptor Signaling Pathway at the Neuromuscular Junction. Journal of Neuroscience, 2000, 20, 8762-8770.	1.7	93
64	Identification of Sites of Incorporation in the Nicotinic Acetylcholine Receptor of a Photoactivatible General Anesthetic. Journal of Biological Chemistry, 2000, 275, 29441-29451.	1.6	70
65	Mapping the Agonist Binding Site of the Nicotinic Acetylcholine Receptor. Journal of Biological Chemistry, 2000, 275, 12651-12660.	1.6	46
66	Probing the Structure of the Nicotinic Acetylcholine Receptor with 4-Benzoylbenzoylcholine, a Novel Photoaffinity Competitive Antagonist. Journal of Biological Chemistry, 2000, 275, 28666-28674.	1.6	26
67	Identification of the Sites of Incorporation of [3H]Ethidium Diazide within the Torpedo Nicotinic Acetylcholine Receptor Ion Channel. Biochemistry, 2000, 39, 11452-11462.	1.2	24
68	Identification of Amino Acids of the Torpedo Nicotinic Acetylcholine Receptor Contributing to the Binding Site for the Noncompetitive Antagonist [³H]Tetracaine. Molecular Pharmacology, 1999, 56, 300-307.	1.0	42
69	Photoaffinity Labeling the Torpedo Nicotinic Acetylcholine Receptor with [³H]Tetracaine, a Nondesensitizing Noncompetitive Antagonist. Molecular Pharmacology, 1999, 56, 290-299.	1.0	52
70	The Steroid Promegestone Is a Noncompetitive Antagonist of theTorpedoNicotinic Acetylcholine Receptor that Interacts with the Lipid-Protein Interface. Molecular Pharmacology, 1999, 55, 269-278.	1.0	58
71	Synthesis and Properties of 3-(2-Hydroxyethyl)-3-n-pentyldiazirine, a Photoactivable General Anesthetic. Journal of Medicinal Chemistry, 1999, 42, 3300-3307.	2.9	43
72	Structure of the Agonist-Binding Sites of the Torpedo Nicotinic Acetylcholine Receptor:â€‰ Affinity-Labeling and Mutational Analyses Identify Î³Tyr-111/Î´Arg-113 as Antagonist Affinity Determinants. Biochemistry, 1999, 38, 6689-6698.	1.2	60

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73	Spatial structure of the M3 transmembrane segment of the nicotinic acetylcholine receptor alpha subunit. FEBS Journal, 1998, 255, 455-461.	0.2	40
74	Identification of tryptophan 55 as the primary site of [3H]nicotine photoincorporation in the Î³-subunit of theTorpedonicotinic acetylcholine receptor. FEBS Letters, 1998, 423, 223-226.	1.3	45
75	Probing the Structure of the Nicotinic Acetylcholine Receptor Ion Channel with the Uncharged Photoactivable Compound [3H]Diazofluorene. Journal of Biological Chemistry, 1998, 273, 8659-8668.	1.6	57
76	Identification of Amino Acids Contributing to High and Low Affinity d-Tubocurarine Sites in the TorpedoNicotinic Acetylcholine Receptor. Journal of Biological Chemistry, 1997, 272, 32940-32950.	1.6	90
77	Active-site peptides of acetylcholinesterase of Electrophorus electricus: labelling of His-440 by 1-bromo-[2-14C]pinacolone and Ser-200 by tritiated diisopropyl fluorophosphate. BBA - Proteins and Proteomics, 1994, 1208, 324-331.	2.1	12
78	Identifying the Lipid-Protein Interface of the Torpedo Nicotinic Acetylcholine Receptor: Secondary Structure Implications. Biochemistry, 1994, 33, 2859-2872.	1.2	221
79	The 87K postsynaptic membrane protein from torpedo is a protein-tyrosine kinase substrate homologous to dystrophin. Neuron, 1993, 10, 511-522.	3.8	152
80	Mapping the lipid-exposed regions in the Torpedo californica nicotinic acetylcholine receptor. Biochemistry, 1992, 31, 3738-3750.	1.2	149
81	Mapping of the acetylcholine binding site of the nicotinic acetylcholine receptor: [3H]nicotine as an agonist photoaffinity label. Biochemistry, 1991, 30, 6987-6997.	1.2	203
82	Variation in the ratio of acetylcholine receptors and the Mr 43,000 receptor-associated protein in embryonic chick myotubes and myoblasts. Developmental Biology, 1990, 140, 437-446.	0.9	9
83	Reactions of 1-bromo-2-[14C]pinacolone with acetylcholinesterase from Torpedo nobiliana. Effects of 5-trimethylammonio-2-pentanone and diisopropyl fluorophosphate. BBA - Proteins and Proteomics, 1989, 997, 167-175.	2.1	18
84	Myristic acid is the NH2-terminal blocking group of the 43-kDa protein ofTorpedonicotinic post-synaptic membranes. FEBS Letters, 1989, 243, 65-69.	1.3	20
85	Photolabeling of membrane-bound Torpedo nicotinic acetylcholine receptor with the hydrophobic probe 3-trifluoromethyl-3-(m-[125I]iodophenyl)diazirine. Biochemistry, 1988, 27, 8741-8751.	1.2	98
86	The 43 kilodalton protein of Torpedo nicotinic postsynaptic membranes: purification and determination of primary structure. Biochemistry, 1987, 26, 7090-7102.	1.2	61
87	Substituted benzenes and phenols as reversible inhibitors of acetylcholinesterase: Polar, trimethyl, and synergistic effects. Bioorganic Chemistry, 1987, 15, 237-249.	2.0	7
88	Desensitization of membrane-bound Torpedo acetylcholine receptor by amine noncompetitive antagonists and aliphatic alcohols: studies of [3H]acetylcholine binding and sodium-22 ion fluxes. Biochemistry, 1984, 23, 4023-4033.	1.2	117
89	Conformations of Torpedo acetylcholine receptor associated with ion transport and desensitization. Biochemistry, 1982, 21, 3460-3467.	1.2	141
90	AGONISTS OF TORPEDO NICOTINIC RECEPTORS: ESSENTIAL ROLE OF A POSITIVE CHARGE. Annals of the New York Academy of Sciences, 1980, 358, 370-373.	1.8	5

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91	Kinetics of binding of [3H]acetylcholine and [3H]carbamoylcholine to Torpedo postsynaptic membranes: slow conformational transitions of the cholinergic receptor. Biochemistry, 1980, 19, 5344-5353.	1.2	144
92	Equilibrium binding of [3H]tubocurarine and [3H]acetylcholine by torpedo postsynaptic membranes: stoichiometry and ligand interactions. Biochemistry, 1979, 18, 5464-5475.	1.2	257
93	Conformational Transitions of the Membrane-Bound Cholinergic Receptor. Jerusalem Symposia on Quantum Chemistry and Biochemistry, 1979, , 293-304.	0.2	2
94	Rotational energy transfer in pure HCN and in HCNâ€rare gas mixtures by microwave double resonance and pressure broadening. Journal of Chemical Physics, 1973, 58, 442-455.	1.2	47
95	Presence of a lattice structure in membrane fragments rich in nicotinic receptor protein from the electric organ of Torpedo marmorata. FEBS Letters, 1973, 33, 109-113.	1.3	203
96	Interaction of a fluorescent ligand with membrane-bound cholinergic receptor from Torpedo marmorata. Biochemistry, 1973, 12, 4855-4864.	1.2	66
97	Microwave double resonance studies of rotational relaxation in polar gases. Journal of Chemical Physics, 1973, 58, 456-467.	1.2	22
98	Purification fromTorpedo marmorataelectric tissue of membrane fragments particularly rich in cholinergic receptor protein. FEBS Letters, 1972, 26, 43-47.	1.3	154