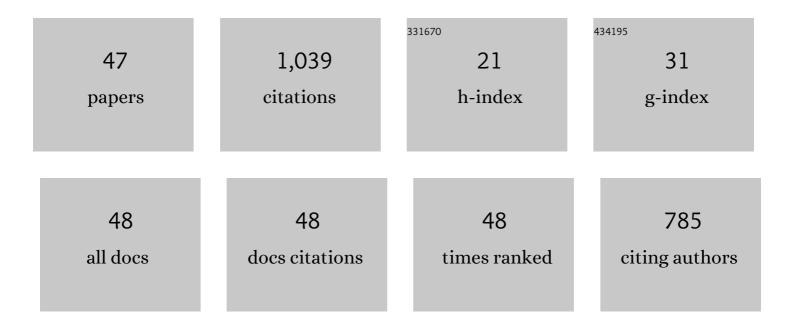
Nicole A Horenstein

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Selective Agonists and Antagonists of α9 Versus α7 Nicotinic Acetylcholine Receptors. ACS Chemical Neuroscience, 2022, 13, 624-637.	3.5	10
2	Comparison of the Anti-inflammatory Properties of Two Nicotinic Acetylcholine Receptor Ligands, Phosphocholine and pCF3-diEPP. Frontiers in Cellular Neuroscience, 2022, 16, 779081.	3.7	11
3	Coffee and cigarettes: Modulation of high and low sensitivity α4β2 nicotinic acetylcholine receptors by n-MP, a biomarker of coffee consumption. Neuropharmacology, 2022, 216, 109173.	4.1	5
4	Therapeutic Targeting of <i>α</i> 7 Nicotinic Acetylcholine Receptors. Pharmacological Reviews, 2021, 73, 1118-1149.	16.0	43
5	Stable desensitization of α7 nicotinic acetylcholine receptors by NS6740 requires interaction with S36 in the orthosteric agonist binding site. European Journal of Pharmacology, 2021, 905, 174179.	3.5	4
6	Comparative genomic analysis of azasugar biosynthesis. AMB Express, 2021, 11, 120.	3.0	5
7	Sulfonium Ligands of the \hat{l} ±7 nAChR. Molecules, 2021, 26, 5643.	3.8	2
8	A silent agonist of α7 nicotinic acetylcholine receptors modulates inflammation ex vivo and attenuates EAE. Brain, Behavior, and Immunity, 2020, 87, 286-300.	4.1	35
9	A Computational Analysis of the Factors Governing the Dynamics of α7 nAChR and Its Homologs. Biophysical Journal, 2020, 119, 1656-1669.	0.5	1
10	Design, synthesis, and electrophysiological evaluation of NS6740 derivatives: Exploration of the structure-activity relationship for alpha7 nicotinic acetylcholine receptor silent activation. European Journal of Medicinal Chemistry, 2020, 205, 112669.	5.5	12
11	Differing Activity Profiles of the Stereoisomers of 2,3,5,6TMP-TQS, a Putative Silent Allosteric Modulator of <i>α</i> 7 nAChR. Molecular Pharmacology, 2020, 98, 292-302.	2.3	12
12	Heteromeric Neuronal Nicotinic Acetylcholine Receptors with Mutant <i>β</i> Subunits Acquire Sensitivity to <i>α</i> 7-Selective Positive Allosteric Modulators. Journal of Pharmacology and Experimental Therapeutics, 2019, 370, 252-268.	2.5	10
13	Synthesis of saccharin-glycoconjugates targeting carbonic anhydrase using a one-pot cyclization/deprotection strategy. Carbohydrate Research, 2019, 476, 65-70.	2.3	8
14	Allosteric Agonism of α7 Nicotinic Acetylcholine Receptors: Receptor Modulation Outside the Orthosteric Site. Molecular Pharmacology, 2019, 95, 606-614.	2.3	24
15	Functional Analysis of a Gene Cluster from <i>Chitinophaga pinensis</i> Involved in Biosynthesis of the Pyrrolidine Azasugar DAB-1. Journal of Natural Products, 2019, 82, 3401-3409.	3.0	6
16	Macroscopic and Microscopic Activation of <i>α</i> 7 Nicotinic Acetylcholine Receptors by the Structurally Unrelated Allosteric Agonist-Positive Allosteric Modulators (ago-PAMs) B-973B and GAT107. Molecular Pharmacology, 2019, 95, 43-61.	2.3	21
17	Cracking the Betel Nut: Cholinergic Activity of Areca Alkaloids and Related Compounds. Nicotine and Tobacco Research, 2019, 21, 805-812.	2.6	25
18	Persistent activation of α7 nicotinic ACh receptors associated with stable induction of different desensitized states. British Journal of Pharmacology, 2018, 175, 1838-1854.	5.4	31

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19	Novel 5-(quinuclidin-3-ylmethyl)-1,2,4-oxadiazoles to investigate the activation of the α7 nicotinic acetylcholine receptor subtype: Synthesis and electrophysiological evaluation. European Journal of Medicinal Chemistry, 2018, 160, 207-228.	5.5	9
20	The Antinociceptive and Anti-Inflammatory Properties of the <i>α</i> 7 nAChR Weak Partial Agonist <i>p</i> -CF ₃ <i>N</i> , <i>N</i> -diethyl- <i>N</i> ′-phenylpiperazine. Journal of Pharmacology and Experimental Therapeutics, 2018, 367, 203-214.	2.5	17
21	Characterization of the PLP-dependent transaminase initiating azasugar biosynthesis. Biochemical Journal, 2018, 475, 2241-2256.	3.7	7
22	Identification of α7 Nicotinic Acetylcholine Receptor Silent Agonists Based on the Spirocyclic Quinuclidineâ€Î" ² â€Isoxazoline Scaffold: Synthesis and Electrophysiological Evaluation. ChemMedChem, 2017, 12, 1335-1348.	3.2	15
23	Anti-inflammatory Silent Agonists. ACS Medicinal Chemistry Letters, 2017, 8, 989-991.	2.8	38
24	Sulfonium as a Surrogate for Ammonium: A New α7 Nicotinic Acetylcholine Receptor Partial Agonist with Desensitizing Activity. Journal of Medicinal Chemistry, 2017, 60, 7928-7934.	6.4	10
25	Experimental and Metabolic Modeling Evidence for a Folate-Cleaving Side-Activity of Ketopantoate Hydroxymethyltransferase (PanB). Frontiers in Microbiology, 2016, 7, 431.	3.5	6
26	Critical Molecular Determinants of α7 Nicotinic Acetylcholine Receptor Allosteric Activation. Journal of Biological Chemistry, 2016, 291, 5049-5067.	3.4	43
27	Dissection of N,N-diethyl-N′-phenylpiperazines as α7 nicotinic receptor silent agonists. Bioorganic and Medicinal Chemistry, 2016, 24, 286-293.	3.0	31
28	Nicotinic Activity of Arecoline, the Psychoactive Element of "Betel Nuts", Suggests a Basis for Habitual Use and Anti-Inflammatory Activity. PLoS ONE, 2015, 10, e0140907.	2.5	96
29	The Minimal Pharmacophore for Silent Agonism of the α7 Nicotinic Acetylcholine Receptor. Journal of Pharmacology and Experimental Therapeutics, 2014, 350, 665-680.	2.5	41
30	The Activity of GAT107, an Allosteric Activator and Positive Modulator of α7 Nicotinic Acetylcholine Receptors (nAChR), Is Regulated by Aromatic Amino Acids That Span the Subunit Interface. Journal of Biological Chemistry, 2014, 289, 4515-4531.	3.4	36
31	Synthesis and evaluation of a conditionally-silent agonist for the α7 nicotinic acetylcholine receptor. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 4145-4149.	2.2	41
32	Point-to-point ligand–receptor interactions across the subunit interface modulate the induction and stabilization of conformational states of alpha7 nAChR by benzylidene anabaseines. Biochemical Pharmacology, 2013, 85, 817-828.	4.4	3
33	Potential State-selective Hydrogen Bond Formation Can Modulate Activation and Desensitization of the α7 Nicotinic Acetylcholine Receptor. Journal of Biological Chemistry, 2012, 287, 21957-21969.	3.4	8
34	Cysteine accessibility analysis of the human alpha7 nicotinic acetylcholine receptor ligand-binding domain identifies L119 as a gatekeeper. Neuropharmacology, 2011, 60, 159-171.	4.1	26
35	Identification of a Gene Cluster that Initiates Azasugar Biosynthesis in <i>Bacillus amyloliquefaciens</i> . ChemBioChem, 2011, 12, 2147-2150.	2.6	30
36	The effective opening of nicotinic acetylcholine receptors with single agonist binding sites. Journal of General Physiology, 2011, 137, 369-384.	1.9	44

NICOLE A HORENSTEIN

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37	Synthesis of 3,5-diazabicyclo [5.1.0] octenes. A new platform to mimic glycosidase transition states. Tetrahedron, 2010, 66, 5566-5572.	1.9	4
38	Tethered Agonist Analogs as Site-Specific Probes for Domains of the Human α7 Nicotinic Acetylcholine Receptor that Differentially Regulate Activation and Desensitization. Molecular Pharmacology, 2010, 78, 1012-1025.	2.3	23
39	Activation and Desensitization of Nicotinic α7-type Acetylcholine Receptors by Benzylidene Anabaseines and Nicotine. Journal of Pharmacology and Experimental Therapeutics, 2009, 329, 791-807.	2.5	83
40	Differential Regulation of Receptor Activation and Agonist Selectivity by Highly Conserved Tryptophans in the Nicotinic Acetylcholine Receptor Binding Site. Journal of Pharmacology and Experimental Therapeutics, 2009, 330, 40-53.	2.5	24
41	Synthesis of H-bonding probes of α7 nAChR agonist selectivity. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 474-476.	2.2	4
42	Multiple Pharmacophores for the Selective Activation of Nicotinic α7-Type Acetylcholine Receptors. Molecular Pharmacology, 2008, 74, 1496-1511.	2.3	52
43	Reversal of Agonist Selectivity by Mutations of Conserved Amino Acids in the Binding Site of Nicotinic Acetylcholine Receptors. Journal of Biological Chemistry, 2007, 282, 5899-5909.	3.4	31
44	A new route into hexahydro-cyclopenta[b]pyrrole-cis-3a,6-diols. Synthesis of constrained bicyclic analogues of pyrrolidine azasugars. Tetrahedron, 2005, 61, 10462-10469.	1.9	13
45	Enzymatic synthesis of [1-14C-N-acetyl, P18O2] cytidine monophosphate neuraminic acid. Journal of Labelled Compounds and Radiopharmaceuticals, 2004, 47, 1007-1017.	1.0	1
46	Synthesis of endo-(3-azabicyclo[3.1.0]hex-6-yl)-methanol and derivatives as new geometric/charge mimics of glycosyltransfer transition states. Tetrahedron Letters, 2004, 45, 9505-9507.	1.4	6
47	Effects at a distance in α7 nAChR selective agonists: benzylidene substitutions that regulate potency and efficacy. Neuropharmacology, 2004, 46, 1023-1038.	4.1	32