

David M Thal

List of Publications by Year in descending order

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32
papers

3,018
citations

304743

22
h-index

526287

27
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35
all docs

35
docs citations

35
times ranked

3942
citing authors

#	ARTICLE	IF	CITATIONS
1	Phase-plate cryo-EM structure of a class B GPCR-G-protein complex. <i>Nature</i> , 2017, 546, 118-123.	27.8	424
2	THE CONCISE GUIDE TO PHARMACOLOGY 2021/22: G protein-coupled receptors. <i>British Journal of Pharmacology</i> , 2021, 178, S27-S156.	5.4	337
3	Structure of the adenosine-bound human adenosine A1 receptor-Gi complex. <i>Nature</i> , 2018, 558, 559-563.	27.8	274
4	Crystal structures of the M1 and M4 muscarinic acetylcholine receptors. <i>Nature</i> , 2016, 531, 335-340.	27.8	272
5	Phase-plate cryo-EM structure of a biased agonist-bound human GLP-1 receptor-Gs complex. <i>Nature</i> , 2018, 555, 121-125.	27.8	263
6	Structural insights into G-protein-coupled receptor allostery. <i>Nature</i> , 2018, 559, 45-53.	27.8	255
7	Structure of the Adenosine A1 Receptor Reveals the Basis for Subtype Selectivity. <i>Cell</i> , 2017, 168, 867-877.e13.	28.9	237
8	Paroxetine Is a Direct Inhibitor of G Protein-Coupled Receptor Kinase 2 and Increases Myocardial Contractility. <i>ACS Chemical Biology</i> , 2012, 7, 1830-1839.	3.4	163
9	Molecular Mechanism of Selectivity among G Protein-Coupled Receptor Kinase 2 Inhibitors. <i>Molecular Pharmacology</i> , 2011, 80, 294-303.	2.3	104
10	Positive allosteric mechanisms of adenosine A1 receptor-mediated analgesia. <i>Nature</i> , 2021, 597, 571-576.	27.8	84
11	Incorporation and Replication of 8-Oxo-deoxyguanosine by the Human Mitochondrial DNA Polymerase. <i>Journal of Biological Chemistry</i> , 2006, 281, 36241-36248.	3.4	76
12	An autoinhibitory helix in the C-terminal region of phospholipase C- β^2 mediates G $\beta\gamma$ activation. <i>Nature Structural and Molecular Biology</i> , 2011, 18, 999-1005.	8.2	71
13	Molecular Determinants of Allosteric Modulation at the M1 Muscarinic Acetylcholine Receptor. <i>Journal of Biological Chemistry</i> , 2014, 289, 6067-6079.	3.4	51
14	Recent advances in the determination of G protein-coupled receptor structures. <i>Current Opinion in Structural Biology</i> , 2018, 51, 28-34.	5.7	51
15	Crystal structure of the M ₅ muscarinic acetylcholine receptor. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 26001-26007.	7.1	48
16	Assembly of High Order G $\beta\gamma$ -Effector Complexes with RGS Proteins. <i>Journal of Biological Chemistry</i> , 2008, 283, 34923-34934.	3.4	46
17	Structures of the human cholecystokinin 1 (CCK1) receptor bound to Gs and Gq mimetic proteins provide insight into mechanisms of G protein selectivity. <i>PLoS Biology</i> , 2021, 19, e3001295.	5.6	41
18	Selective G protein signaling driven by substance P-neurokinin receptor dynamics. <i>Nature Chemical Biology</i> , 2022, 18, 109-115.	8.0	40

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19	Toward an understanding of the structural basis of allostery in muscarinic acetylcholine receptors. <i>Journal of General Physiology</i> , 2018, 150, 1360-1372.	1.9	38
20	Unique Hydrophobic Extension of the RGS2 Amphipathic Helix Domain Imparts Increased Plasma Membrane Binding and Function Relative to Other RGS R4/B Subfamily Members. <i>Journal of Biological Chemistry</i> , 2007, 282, 33064-33075.	3.4	32
21	Clickable Photoaffinity Ligands for Metabotropic Glutamate Receptor 5 Based on Select Acetylenic Negative Allosteric Modulators. <i>ACS Chemical Biology</i> , 2016, 11, 1870-1879.	3.4	26
22	The structural determinants of the bitopic binding mode of a negative allosteric modulator of the dopamine D2 receptor. <i>Biochemical Pharmacology</i> , 2018, 148, 315-328.	4.4	26
23	Cluster cytometry for high-capacity bioanalysis. <i>Cytometry Part A: the Journal of the International Society for Analytical Cytology</i> , 2012, 81A, 419-429.	1.5	20
24	The action of a negative allosteric modulator at the dopamine D2 receptor is dependent upon sodium ions. <i>Scientific Reports</i> , 2018, 8, 1208.	3.3	16
25	Identification of a Novel Allosteric Site at the M5 Muscarinic Acetylcholine Receptor. <i>ACS Chemical Neuroscience</i> , 2021, 12, 3112-3123.	3.5	6
26	Biased Profile of Xanomeline at the Recombinant Human M ₄ Muscarinic Acetylcholine Receptor. <i>ACS Chemical Neuroscience</i> , 2022, 13, 1206-1218.	3.5	6
27	The P2X1 receptor as a therapeutic target. <i>Purinergic Signalling</i> , 2022, 18, 421-433.	2.2	6
28	Acetylcholine receptors (muscarinic) in GtoPdb v.2021.2. <i>IUPHAR/BPS Guide To Pharmacology CITE</i> , 2021, .	0.2	0
29	Acetylcholine receptors (muscarinic) in GtoPdb v.2021.3. <i>IUPHAR/BPS Guide To Pharmacology CITE</i> , 2021, .	0.2	0
30	How G α q Regulates PIP 2 Hydrolysis: Molecular Mechanisms and Prospects for Drug Development. <i>FASEB Journal</i> , 2012, 26, 667.1.	0.5	0
31	New Therapeutics Targeting Heart Failure: Development of GRK2 Selective Inhibitors. <i>FASEB Journal</i> , 2012, 26, 665.8.	0.5	0
32	P598. Exploring the Molecular Determinants for Functional Selectivity of the Antipsychotic Xanomeline at Muscarinic Acetylcholine Receptors. <i>Biological Psychiatry</i> , 2022, 91, S331.	1.3	0