

# Mark J Henderson

## List of Publications by Year in descending order

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Version: 2024-02-01

32  
papers

1,186  
citations

430874

18  
h-index

414414

32  
g-index

33  
all docs

33  
docs citations

33  
times ranked

2305  
citing authors

#	ARTICLE	IF	CITATIONS
1	Discovery of Small-Molecule VapC1 Nuclease Inhibitors by Virtual Screening and Scaffold Hopping from an Atomic Structure Revealing Protein-Protein Interactions with a Native VapB1 Inhibitor. <i>Journal of Chemical Information and Modeling</i> , 2022, 62, 1249-1258.	5.4	3
2	High-Throughput Cellular Thermal Shift Assay Using Acoustic Transfer of Protein Lysates. <i>ACS Chemical Biology</i> , 2022, , .	3.4	8
3	Discovery and Optimization of Pyrrolopyrimidine Derivatives as Selective Disruptors of the Perinucleolar Compartment, a Marker of Tumor Progression toward Metastasis. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 8303-8331.	6.4	4
4	The SARS-CoV-2 Cytopathic Effect Is Blocked by Lysosome Alkalinizing Small Molecules. <i>ACS Infectious Diseases</i> , 2021, 7, 1389-1408.	3.8	74
5	High-Throughput Detection of Ligand-Protein Binding Using a SplitLuc Cellular Thermal Shift Assay. <i>Methods in Molecular Biology</i> , 2021, 2365, 21-41.	0.9	7
6	Discovery and Optimization of 2-Hydroxy-2-Pyridin-2-one Inhibitors of Mutant Isocitrate Dehydrogenase 1 for the Treatment of Cancer. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 4913-4946.	6.4	12
7	A target-agnostic screen identifies approved drugs to stabilize the endoplasmic reticulum-resident proteome. <i>Cell Reports</i> , 2021, 35, 109040.	6.4	18
8	The AKT modulator A-443654 reduces $\alpha$ -synuclein expression and normalizes ER stress and autophagy. <i>Journal of Biological Chemistry</i> , 2021, 297, 101191.	3.4	7
9	High-Throughput Cellular Thermal Shift Assays in Research and Drug Discovery. <i>SLAS Discovery</i> , 2020, 25, 137-147.	2.7	35
10	A Comparative Study of Target Engagement Assays for HDAC1 Inhibitor Profiling. <i>SLAS Discovery</i> , 2020, 25, 253-264.	2.7	9
11	Pyrazole-Based Lactate Dehydrogenase Inhibitors with Optimized Cell Activity and Pharmacokinetic Properties. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 10984-11011.	6.4	30
12	Design, Synthesis, and Biological Evaluation of Quinazolin-4-one-Based Hydroxamic Acids as Dual PI3K/HDAC Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 4256-4292.	6.4	59
13	Compound screening in cell-based models of tau inclusion formation: Comparison of primary neuron and HEK293 cell assays. <i>Journal of Biological Chemistry</i> , 2020, 295, 4001-4013.	3.4	10
14	A small molecule inhibitor of ER-to-cytosol protein dislocation exhibits anti-dengue and anti-Zika virus activity. <i>Scientific Reports</i> , 2019, 9, 10901.	3.3	15
15	Physiologically relevant orthogonal assays for the discovery of small-molecule modulators of WIP1 phosphatase in high-throughput screens. <i>Journal of Biological Chemistry</i> , 2019, 294, 17654-17668.	3.4	6
16	Discovery of endoplasmic reticulum calcium stabilizers to rescue ER-stressed podocytes in nephrotic syndrome. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 14154-14163.	7.1	39
17	Ipomoeassin F Binds Sec61 to Inhibit Protein Translocation. <i>Journal of the American Chemical Society</i> , 2019, 141, 8450-8461.	13.7	58
18	Insights into the Action of Inhibitor Enantiomers against Histone Lysine Demethylase 5A. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 3193-3208.	6.4	9

#	ARTICLE	IF	CITATIONS
19	KDEL Receptors Are Differentially Regulated to Maintain the ER Proteome under Calcium Deficiency. <i>Cell Reports</i> , 2018, 25, 1829-1840.e6.	6.4	93
20	Canvass: A Crowd-Sourced, Natural-Product Screening Library for Exploring Biological Space. <i>ACS Central Science</i> , 2018, 4, 1727-1741.	11.3	32
21	Structure-Based Engineering of Irreversible Inhibitors against Histone Lysine Demethylase KDM5A. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 10588-10601.	6.4	28
22	High-throughput screening with nucleosome substrate identifies small-molecule inhibitors of the human histone lysine methyltransferase NSD2. <i>Journal of Biological Chemistry</i> , 2018, 293, 13750-13765.	3.4	46
23	KDM5 histone demethylases repress immune response via suppression of STING. <i>PLoS Biology</i> , 2018, 16, e2006134.	5.6	106
24	A widely-applicable high-throughput cellular thermal shift assay (CETSA) using split Nano Luciferase. <i>Scientific Reports</i> , 2018, 8, 9472.	3.3	65
25	Assessing inhibitors of mutant isocitrate dehydrogenase using a suite of pre-clinical discovery assays. <i>Scientific Reports</i> , 2017, 7, 12758.	3.3	59
26	Connecting Neuronal Cell Protective Pathways and Drug Combinations in a Huntington's Disease Model through the Application of Quantitative Systems Pharmacology. <i>Scientific Reports</i> , 2017, 7, 17803.	3.3	22
27	Longitudinal monitoring of Gaussia and Nano luciferase activities to concurrently assess ER calcium homeostasis and ER stress in vivo. <i>PLoS ONE</i> , 2017, 12, e0175481.	2.5	11
28	Structural Basis for KDM5A Histone Lysine Demethylase Inhibition by Diverse Compounds. <i>Cell Chemical Biology</i> , 2016, 23, 769-781.	5.2	80
29	Monitoring Endoplasmic Reticulum Calcium Homeostasis Using a <i>Gaussia</i> Luciferase SERCaMP. <i>Journal of Visualized Experiments</i> , 2015, , .	0.3	9
30	A Low Affinity GCaMP3 Variant (GCaMPer) for Imaging the Endoplasmic Reticulum Calcium Store. <i>PLoS ONE</i> , 2015, 10, e0139273.	2.5	51
31	SERCaMP: a carboxy-terminal protein modification that enables monitoring of ER calcium homeostasis. <i>Molecular Biology of the Cell</i> , 2014, 25, 2828-2839.	2.1	54
32	Mesencephalic Astrocyte-derived Neurotrophic Factor (MANF) Secretion and Cell Surface Binding Are Modulated by KDEL Receptors. <i>Journal of Biological Chemistry</i> , 2013, 288, 4209-4225.	3.4	127