

Armand B Cagnetta Iii

List of Publications by Year in descending order

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Version: 2024-02-01

18
papers

1,721
citations

430442

18
h-index

839053

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19
docs citations

19
times ranked

2799
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification of cell wall synthesis inhibitors active against <i>Mycobacterium tuberculosis</i> by competitive activity-based protein profiling. <i>Cell Chemical Biology</i> , 2022, 29, 883-896.e5.	2.5	20
2	Notum produced by Paneth cells attenuates regeneration of aged intestinal epithelium. <i>Nature</i> , 2019, 571, 398-402.	13.7	166
3	Global Portrait of Protein Targets of Metabolites of the Neurotoxic Compound BIA 10-2474. <i>ACS Chemical Biology</i> , 2019, 14, 192-197.	1.6	40
4	Pharmacological convergence reveals a lipid pathway that regulates <i>C. elegans</i> lifespan. <i>Nature Chemical Biology</i> , 2019, 15, 453-462.	3.9	35
5	Selective Irreversible Inhibitors of the Wnt-Deacylating Enzyme NOTUM Developed by Activity-Based Protein Profiling. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 563-568.	1.3	39
6	A Screen for Protein-Protein Interactions in Live <i>Mycobacteria</i> Reveals a Functional Link between the Virulence-Associated Lipid Transporter LprG and the Mycolyltransferase Antigen 85A. <i>ACS Infectious Diseases</i> , 2017, 3, 336-348.	1.8	23
7	Design of Benzothiazin-3-one 1,1-Dioxides as a New Class of Irreversible Serine Hydrolase Inhibitors: Discovery of a Uniquely Selective PNPLA4 Inhibitor. <i>Journal of the American Chemical Society</i> , 2017, 139, 7052-7061.	6.6	25
8	Activity-based protein profiling reveals off-target proteins of the FAAH inhibitor BIA 10-2474. <i>Science</i> , 2017, 356, 1084-1087.	6.0	251
9	Multicomponent mapping of boron chemotypes furnishes selective enzyme inhibitors. <i>Nature Communications</i> , 2017, 8, 1760.	5.8	30
10	AIG1 and ADTRP are atypical integral membrane hydrolases that degrade bioactive FAHFAs. <i>Nature Chemical Biology</i> , 2016, 12, 367-372.	3.9	62
11	A Global Map of Lipid-Binding Proteins and Their Ligandability in Cells. <i>Cell</i> , 2015, 161, 1668-1680.	13.5	188
12	Selective N-Hydroxyhydantoin Carbamate Inhibitors of Mammalian Serine Hydrolases. <i>Chemistry and Biology</i> , 2015, 22, 928-937.	6.2	52
13	Facile synthesis of borofragments and their evaluation in activity-based protein profiling. <i>Chemical Communications</i> , 2015, 51, 3608-3611.	2.2	25
14	Remodeling Natural Products: Chemistry and Serine Hydrolase Activity of a Rocaglate-Derived β -Lactone. <i>Journal of the American Chemical Society</i> , 2014, 136, 2659-2664.	6.6	37
15	Proteome-Wide Reactivity Profiling Identifies Diverse Carbamate Chemotypes Tuned for Serine Hydrolase Inhibition. <i>ACS Chemical Biology</i> , 2013, 8, 1590-1599.	1.6	105
16	Proteome-wide mapping of cholesterol-interacting proteins in mammalian cells. <i>Nature Methods</i> , 2013, 10, 259-264.	9.0	350
17	Evaluation of NHS Carbamates as a Potent and Selective Class of Endocannabinoid Hydrolase Inhibitors. <i>ACS Chemical Neuroscience</i> , 2013, 4, 1322-1332.	1.7	116
18	Highly Selective Inhibitors of Monoacylglycerol Lipase Bearing a Reactive Group that Is Bioisosteric with Endocannabinoid Substrates. <i>Chemistry and Biology</i> , 2012, 19, 579-588.	6.2	155