

Rachel L Palte

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/548505/publications.pdf>

Version: 2024-02-01

8
papers

111
citations

1307594

7
h-index

1588992

8
g-index

9
all docs

9
docs citations

9
times ranked

183
citing authors

#	ARTICLE	IF	CITATIONS
1	The Discovery of Two Novel Classes of 5,5-Bicyclic Nucleoside-Derived PRMT5 Inhibitors for the Treatment of Cancer. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 3911-3939.	6.4	16
2	Development of a Flexible and Robust Synthesis of Tetrahydrofuro[3,4- <i>b</i>]furan Nucleoside Analogues. <i>Journal of Organic Chemistry</i> , 2021, 86, 5142-5151.	3.2	7
3	Cryo-EM structures of inhibitory antibodies complexed with arginase 1 provide insight into mechanism of action. <i>Communications Biology</i> , 2021, 4, 927.	4.4	2
4	Structure-Based Discovery of Proline-Derived Arginase Inhibitors with Improved Oral Bioavailability for Immuno-Oncology. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 1380-1388.	2.8	11
5	Comprehensive Strategies to Bicyclic Prolines: Applications in the Synthesis of Potent Arginase Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 1678-1688.	2.8	9
6	Discovery of <i>N</i> -(Indazol-3-yl)piperidine-4-carboxylic Acids as ROR α Allosteric Inhibitors for Autoimmune Diseases. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 114-119.	2.8	18
7	Allosteric Modulation of Protein Arginine Methyltransferase 5 (PRMT5). <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 1688-1693.	2.8	30
8	Discovery and Optimization of Rationally Designed Bicyclic Inhibitors of Human Arginase to Enhance Cancer Immunotherapy. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 582-588.	2.8	18