

Luyi Huang

List of Publications by Year in descending order

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Version: 2024-02-01

11
papers

545
citations

1163117

8
h-index

1372567

10
g-index

11
all docs

11
docs citations

11
times ranked

920
citing authors

#	ARTICLE	IF	CITATIONS
1	Dopamine-loaded blood exosomes targeted to brain for better treatment of Parkinson's disease. <i>Journal of Controlled Release</i> , 2018, 287, 156-166.	9.9	329
2	Porphyrins with intense absorptivity: highly efficient sensitizers with a photovoltaic efficiency of up to 10.7% without a cosensitizer and a coabsorbate. <i>Journal of Materials Chemistry A</i> , 2016, 4, 11829-11834.	10.3	56
3	Discovery of New SIRT2 Inhibitors by Utilizing a Consensus Docking/Scoring Strategy and Structure-Activity Relationship Analysis. <i>Journal of Chemical Information and Modeling</i> , 2017, 57, 669-679.	5.4	33
4	Identification of 5-(2,3-Dihydro-1 <i>H</i> -indol-5-yl)-7 <i>H</i> -pyrrolo[2,3- <i>d</i>]pyrimidin-4-amine Derivatives as a New Class of Receptor-Interacting Protein Kinase 1 (RIPK1) Inhibitors, Which Showed Potent Activity in a Tumor Metastasis Model. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 11398-11414.	6.4	33
5	Design, synthesis and biological evaluation of a novel tubulin inhibitor 7a3 targeting the colchicine binding site. <i>European Journal of Medicinal Chemistry</i> , 2018, 156, 162-179.	5.5	28
6	Discovery and structure-activity relationship studies of 1-aryl-1 <i>H</i> -naphtho[2,3- <i>d</i>][1,2,3]triazole-4,9-dione derivatives as potent dual inhibitors of indoleamine 2,3-dioxygenase 1 (IDO1) and tryptophan 2,3-dioxygenase (TDO). <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112703.	5.5	22
7	Structure-Guided Discovery of a Potent and Selective Cell-Active Inhibitor of SETDB1 Tudor Domain. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 8760-8765.	13.8	20
8	Discovery of Pyrrolo[3,2- <i>d</i>]pyrimidin-4-one Derivatives as a New Class of Potent and Cell-Active Inhibitors of P300/CBP-Associated Factor Bromodomain. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 4526-4542.	6.4	11
9	Discovery of 5-(5-fluoro-1 <i>H</i> -pyrrolo[2,3- <i>b</i>]pyridin-3-yl)pyrazin-2(1 <i>H</i>)-one derivatives as new potent PB2 inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 1609-1613.	2.2	7
10	Structure-Based Discovery of N-Sulfonylpiperidine-3-Carboxamides as Novel Capsid Assembly Modulators for Potent Inhibition of HBV Replication. <i>Viruses</i> , 2022, 14, 348.	3.3	5
11	Structure-Guided Discovery of a Potent and Selective Cell-Active Inhibitor of SETDB1 Tudor Domain. <i>Angewandte Chemie</i> , 2021, 133, 8842-8847.	2.0	1