

Phillip P Sharp

List of Publications by Year in descending order

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Version: 2024-02-01

20
papers

4,683
citations

623734

14
h-index

794594

19
g-index

23
all docs

23
docs citations

23
times ranked

11683
citing authors

#	ARTICLE	IF	CITATIONS
1	Inhibition of the Sec61 translocon overcomes cytokine-induced glucocorticoid resistance in T-cell acute lymphoblastic leukaemia. <i>British Journal of Haematology</i> , 2022, , .	2.5	6
2	Inhibitors of Eukaryotic Translational Machinery as Therapeutic Agents. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 2436-2465.	6.4	13
3	Total Syntheses of the 3 <i>H</i> -Pyrrolo[2,3- <i>c</i>]quinolone-Containing Alkaloids Marinoquinolines A, F, K, and Aplidiopsamine A Using a Palladium-Catalyzed Ullmann Cross-Coupling/Reductive Cyclization Pathway. <i>Journal of Organic Chemistry</i> , 2020, 85, 650-663.	3.2	14
4	A SARS-CoV-2 protein interaction map reveals targets for drug repurposing. <i>Nature</i> , 2020, 583, 459-468.	27.8	3,542
5	A small molecule interacts with VDAC2 to block mouse BAK-driven apoptosis. <i>Nature Chemical Biology</i> , 2019, 15, 1057-1066.	8.0	30
6	Protein Translocation Inhibitors Overcome Cytokine-Induced Glucocorticoid Resistance in T-Cell Acute Lymphoblastic Leukemia. <i>Blood</i> , 2019, 134, 805-805.	1.4	0
7	Comparative Flavivirus-Host Protein Interaction Mapping Reveals Mechanisms of Dengue and Zika Virus Pathogenesis. <i>Cell</i> , 2018, 175, 1931-1945.e18.	28.9	252
8	Mechanistic Studies on the Base-Promoted Conversion of Alkoxy-Substituted, Ring-Fused <i>gem</i> -Dihalocyclopropanes into Furans: Evidence for a Process Involving Electrocyclic Ring Closure of a Carbonyl Ylide Intermediate. <i>Journal of Organic Chemistry</i> , 2018, 83, 13678-13690.	3.2	7
9	Synthesis of a GlcNAcylated arginine building block for the solid phase synthesis of death domain glycopeptide fragments. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 2895-2900.	3.0	8
10	Synthesis of rhamnosylated arginine glycopeptides and determination of the glycosidic linkage in bacterial elongation factor P. <i>Chemical Science</i> , 2017, 8, 2296-2302.	7.4	23
11	Design, Synthesis, and Biological Activity of 1,2,3-Triazolobenzodiazepine BET Bromodomain Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2017, 8, 1298-1303.	2.8	23
12	CIS is a potent checkpoint in NK cell-mediated tumor immunity. <i>Nature Immunology</i> , 2016, 17, 816-824.	14.5	289
13	BET inhibition represses miR17-92 to drive BIM-initiated apoptosis of normal and transformed hematopoietic cells. <i>Leukemia</i> , 2016, 30, 1531-1541.	7.2	29
14	BET inhibitors induce apoptosis through a MYC independent mechanism and synergise with CDK inhibitors to kill osteosarcoma cells. <i>Scientific Reports</i> , 2015, 5, 10120.	3.3	103
15	A RIPK2 inhibitor delays NOD signalling events yet prevents inflammatory cytokine production. <i>Nature Communications</i> , 2015, 6, 6442.	12.8	112
16	Simplified Silvestrol Analogues with Potent Cytotoxic Activity. <i>ChemMedChem</i> , 2014, 9, 1556-1566.	3.2	16
17	Evaluation of functional groups as acetyl-lysine mimetics for BET bromodomain inhibition. <i>MedChemComm</i> , 2014, 5, 1834-1842.	3.4	24
18	BET bromodomain inhibitors: a patent review. <i>Expert Opinion on Therapeutic Patents</i> , 2014, 24, 185-199.	5.0	104

#	ARTICLE	IF	CITATIONS
19	Consecutive Gold(I)-Catalyzed Cyclization Reactions of <i>o</i> -(Buta-1,3-dien-1-yl)-Substituted <i>N</i> -Aryl Ureas: A One-Pot Synthesis of Pyrimido[1,6- <i>a</i>]indol-1(2 <i>H</i>)-ones and Related Systems. <i>Organic Letters</i> , 2013, 15, 2616-2619.	4.6	51
20	Dichlorocarbene adducts of alkyl enol ethers as precursors to furans: application to a total synthesis of the furanosesquiterpene (±)-palescensin A. <i>Tetrahedron Letters</i> , 2006, 47, 6817-6820.	1.4	30