Matthias Gehringer

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41 982 5.6 ext. papers ext. citations avg, IF 5.04

L-index

#	Paper	IF	Citations
31	Emerging and Re-Emerging Warheads for Targeted Covalent Inhibitors: Applications in Medicinal Chemistry and Chemical Biology. <i>Journal of Medicinal Chemistry</i> , 2019 , 62, 5673-5724	8.3	218
30	Inhibitors of c-Jun N-terminal kinases: an update. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 72-95	8.3	63
29	Selective JAK3 Inhibitors with a Covalent Reversible Binding Mode Targeting a New Induced Fit Binding Pocket. <i>Cell Chemical Biology</i> , 2016 , 23, 1335-1340	8.2	62
28	The Macrolide Toxin Mycolactone Promotes Bim-Dependent Apoptosis in Buruli Ulcer through Inhibition of mTOR. <i>ACS Chemical Biology</i> , 2017 , 12, 1297-1307	4.9	52
27	Silica-supported l-proline organocatalysts for asymmetric aldolisation. <i>Tetrahedron: Asymmetry</i> , 2009 , 20, 2880-2885		44
26	c-Jun N-terminal kinase inhibitors: a patent review (2010 - 2014). <i>Expert Opinion on Therapeutic Patents</i> , 2015 , 25, 849-72	6.8	39
25	Design and synthesis of tricyclic JAK3 inhibitors with picomolar affinities as novel molecular probes. <i>ChemMedChem</i> , 2014 , 9, 277-81	3.7	29
24	Development, Optimization, and Structure-Activity Relationships of Covalent-Reversible JAK3 Inhibitors Based on a Tricyclic Imidazo[5,4-d]pyrrolo[2,3-b]pyridine Scaffold. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 5350-5366	8.3	27
23	Novel hinge-binding motifs for Janus kinase 3 inhibitors: a comprehensive structure-activity relationship study on tofacitinib bioisosteres. <i>ChemMedChem</i> , 2014 , 9, 2516-27	3.7	25
22	Recent advances in JAK3 inhibition: Isoform selectivity by covalent cysteine targeting. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017 , 27, 4229-4237	2.9	23
21	The chemistry and biology of mycolactones. Beilstein Journal of Organic Chemistry, 2017, 13, 1596-1660	2.5	23
20	Click Chemistry: Novel Applications in Cell Biology and Drug Discovery. <i>Angewandte Chemie - International Edition</i> , 2017 , 56, 15504-15505	16.4	22
19	Developing Small-Molecule Inhibitors of HECT-Type Ubiquitin Ligases for Therapeutic Applications: Challenges and Opportunities. <i>ChemBioChem</i> , 2018 , 19, 2123-2135	3.8	21
18	Solution-phase parallel synthesis of ruxolitinib-derived Janus kinase inhibitors via copper-catalyzed azide-alkyne cycloaddition. <i>ACS Combinatorial Science</i> , 2015 , 17, 5-10	3.9	14
17	A direct enzyme-linked immunosorbent assay (ELISA) for the quantitative evaluation of Janus Kinase 3 (JAK3) inhibitors. <i>Analytical Methods</i> , 2014 , 6, 8817-8822	3.2	11
16	New insights into novel inhibitors against deoxyhypusine hydroxylase from plasmodium falciparum: compounds with an iron chelating potential. <i>Amino Acids</i> , 2015 , 47, 1155-66	3.5	9
15	Discovery of a Novel Class of Covalent Dual Inhibitors Targeting the Protein Kinases BMX and BTK. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	6

LIST OF PUBLICATIONS

14	Tofacitinib and analogs as inhibitors of the histone kinase PRK1 (PKN1). <i>Future Medicinal Chemistry</i> , 2016 , 8, 1537-51	4.1	6
13	Development of an ELISA for the quantification of mycolactone, the cytotoxic macrolide toxin of Mycobacterium ulcerans. <i>PLoS Neglected Tropical Diseases</i> , 2020 , 14, e0008357	4.8	5
12	Configurationally Stabilized Analogs of Exotoxins Mycolactones A and B Reveal the Importance of Side Chain Geometry for Mycolactone Virulence. <i>Organic Letters</i> , 2019 , 21, 5853-5857	6.2	5
11	Current jakinibs for the treatment of rheumatoid arthritis: a systematic review. Inflammopharmacology, 2021 , 29, 595-615	5.1	5
10	Covalent Kinase Inhibitors: An Overview. <i>Topics in Medicinal Chemistry</i> , 2020 , 43-94	0.4	3
9	Nocathioamides, Uncovered by a Tunable Metabologenomic Approach, Define a Novel Class of Chimeric Lanthipeptides. <i>Angewandte Chemie - International Edition</i> , 2021 , 60, 16472-16479	16.4	2
8	Covalent Janus Kinase 3 Inhibitors. <i>Topics in Medicinal Chemistry</i> , 2020 , 225-256	0.4	1
7	An Antigen Capture Assay for the Detection of Mycolactone, the Polyketide Toxin of. <i>Journal of Immunology</i> , 2021 , 206, 2753-2762	5.3	1
6	Chemical Probes for Understudied Kinases: Challenges and Opportunities. <i>Journal of Medicinal Chemistry</i> , 2021 ,	8.3	1
5	N-(6-Chloro-3-nitropyridin-2-yl)-5-(1-methyl-1H-pyrazol-4-yl)isoquinolin-3-amine. <i>MolBank</i> , 2021 , 2021, M1181	0.5	O
4	Gefitinib-Tamoxifen Hybrid Ligands as Potent Agents against Triple-Negative Breast Cancer Journal of Medicinal Chemistry, 2022 , 65, 4616-4632	8.3	О
3	Metabolism of a novel skepinone L-like p38 mitogen-activated protein kinase inhibitor. <i>MedChemComm</i> , 2014 , 5, 808	5	
2	2-(3-{(3R,4R)-4-Methyl-3-[meth-yl(7H-pyrrolo-[2,3-d]pyrimidin-4-yl)amino]-piperidin-1-yl}oxetan-3-yl)ac monohydrate. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2014 , 70, o382-3	eto-nit	rile
1	tert-Butyl N-[(3R,4R)-1-(2-cyano-acet-yl)-4-methyl-piperidin-3-yl]-N-methyl-carbamate. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2013 , 69, 0935		