Alan M Jones

List of Publications by Year in descending order

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Version: 2024-02-01

471061 580395 25 49 798 17 h-index citations g-index papers 52 52 52 1072 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	The Shono-type electroorganic oxidation of unfunctionalised amides. Carbon–carbon bond formation via electrogenerated <i>N</i> -acyliminium ions. Beilstein Journal of Organic Chemistry, 2014, 10, 3056-3072.	1.3	91
2	A novel role for small molecule glycomimetics in the protection against lipid-induced endothelial dysfunction: Involvement of Akt/eNOS and Nrf2/ARE signaling. Biochimica Et Biophysica Acta - General Subjects, 2017, 1861, 3311-3322.	1.1	58
3	Structure-based design, discovery and development of checkpoint kinase inhibitors as potential anticancer therapies. Expert Opinion on Drug Discovery, 2013, 8, 621-640.	2.5	57
4	Thiazolidine derivatives as potent and selective inhibitors of the PIM kinase family. Bioorganic and Medicinal Chemistry, 2017, 25, 2657-2665.	1.4	40
5	Identification of different side effects between PARP inhibitors and their polypharmacological multiâ€ŧarget rationale. British Journal of Clinical Pharmacology, 2022, 88, 742-752.	1.1	38
6	Exploiting Protein Conformational Change to Optimize Adenosine-Derived Inhibitors of HSP70. Journal of Medicinal Chemistry, 2016, 59, 4625-4636.	2.9	29
7	Structure–activity relationships and colorimetric properties of specific probes for the putative cancer biomarker human arylamine N-acetyltransferase 1. Bioorganic and Medicinal Chemistry, 2014, 22, 3030-3054.	1.4	28
8	Diversity-Oriented Synthetic Strategies Applied to Cancer Chemical Biology and Drug Discovery. Molecules, 2014, 19, 17221-17255.	1.7	27
9	An experimentalist's guide to electrosynthesis: the Shono oxidation. Tetrahedron Letters, 2015, 56, 6863-6867.	0.7	24
10	A fragment-based approach applied to a highly flexible target: Insights and challenges towards the inhibition of HSP70 isoforms. Scientific Reports, 2016, 6, 34701.	1.6	24
11	Binding to an Unusual Inactive Kinase Conformation by Highly Selective Inhibitors of Inositol-Requiring Enzyme 1α Kinase-Endoribonuclease. Journal of Medicinal Chemistry, 2019, 62, 2447-2465.	2.9	23
12	Medical errors: Healthcare professionals' perspective at a tertiary hospital in Kuwait. PLoS ONE, 2019, 14, e0217023.	1.1	22
13	Sulfation made simple: a strategy for synthesising sulfated molecules. Chemical Communications, 2019, 55, 4319-4322.	2.2	22
14	Organoruthenium Complexes with Benzo-Fused Pyrithiones Overcome Platinum Resistance in Ovarian Cancer Cells. Cancers, 2021, 13, 2493.	1.7	22
15	Clinical Potential of Targeting Fibroblast Growth Factorâ€23 and αKlotho in the Treatment of Uremic Cardiomyopathy. Journal of the American Heart Association, 2020, 9, e016041.	1.6	20
16	Diabetic endothelial colony forming cells have the potential for restoration with glycomimetics. Scientific Reports, 2019, 9, 2309.	1.6	19
17	Metabolism Mimicry: An Electrosynthetic Method for the Selective Deethylation of Tertiary Benzamides. ChemElectroChem, 2019, 6, 4284-4291.	1.7	17
18	Lysyl Oxidase Likeâ€⊋ (LOXL2): An Emerging Oncology Target. Advanced Therapeutics, 2020, 3, 1900119.	1.6	17

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19	Selective C–H bond electro-oxidation of benzylic acetates and alcohols to benzaldehydes. Organic and Biomolecular Chemistry, 2017, 15, 10010-10015.	1.5	15
20	Synthesis and Spectroscopic Analysis of Piperine- and Piperlongumine-Inspired Natural Product Scaffolds and Their Molecular Docking with IL-1Î ² and NF-Î ² B Proteins. Molecules, 2020, 25, 2841.	1.7	15
21	Metabolismâ€Inspired Electrosynthesis. ChemElectroChem, 2019, 6, 4093-4104.	1.7	14
22	Relevance of physicochemical properties and functional pharmacology data to predict the clinical safety profile of direct oral anticoagulants. Pharmacology Research and Perspectives, 2020, 8, e00603.	1.1	14
23	An expedient synthesis of oxazepino and oxazocino quinazolines. Tetrahedron Letters, 2015, 56, 6478-6483.	0.7	13
24	Electrically Driven <i>N</i> (_{sp} ²)â€" <i>C</i> (_{sp} ^{2/3}) Bond Cleavage of Sulfonamides. ACS Sustainable Chemistry and Engineering, 2020, 8, 3487-3493.	3.2	13
25	Small Molecule Glycomimetics Inhibit Vascular Calcification via c-Met/Notch3/HES1 Signalling. Cellular Physiology and Biochemistry, 2019, 53, 323-336.	1.1	13
26	Parallel synthesis and spectroscopic analysis of a collection of heterocycles based on the diazabenz[e]aceanthrylene core structure. Tetrahedron, 2009, 65, 563-578.	1.0	11
27	Asymmetric Catalytic Oxidative Cleavage of Polycyclic Systems: The Synthesis of Atropisomeric Diazonanes and Diazecanes. Chemistry - A European Journal, 2011, 17, 5714-5718.	1.7	10
28	A core switching strategy to pyrrolo[2,3-b]quinolines and diazocino[1,2-a]indolinones. Organic and Biomolecular Chemistry, 2016, 14, 8998-9011.	1.5	8
29	A Mitsunobu reaction to functionalized cyclic and bicyclic N-arylamines. Tetrahedron Letters, 2018, 59, 238-242.	0.7	8
30	Chemical Methods for N―and Oâ€Sulfation of Small Molecules, Amino Acids and Peptides. ChemBioChem, 2020, 21, 938-942.	1.3	8
31	A novel exchange method to access sulfated molecules. Scientific Reports, 2020, 10, 16559.	1.6	8
32	Diallingâ€In New Reactivity into the Shonoâ€type Anodic Oxidation Reaction. Chemical Record, 2020, 21, 2120-2129.	2.9	8
33	The modulatory role of sulfated and non-sulfated small molecule heparan sulfate-glycomimetics in endothelial dysfunction: absolute structural clarification, molecular docking and simulated dynamics, SAR analyses and ADMET studies. RSC Medicinal Chemistry, 2021, 12, 779-790.	1.7	8
34	Synthesis and characterisation of medium-sized ring systems by oxidative cleavage. Part 2: Insights from the study of ring expanded analogues. Tetrahedron, 2010, 66, 9694-9702.	1.0	7
35	The chemical reactivity of a known anti-psoriasis drug. Part 1: Further insights into the products resulting from oxidative cleavage. Tetrahedron, 2010, 66, 9667-9674.	1.0	7
36	Total synthesis and structural revision of a mangrove alkaloid. RSC Advances, 2017, 7, 48754-48758.	1.7	7

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37	A Structureâ€Reactivity Relationship of the Tandem Asymmetric Dihydroxylation on a Biologically Relevant Diene: Influence of Remote Stereocenters on Diastereofacial Selectivity. European Journal of Organic Chemistry, 2019, 2019, 7568-7577.	1.2	7
38	Polypharmacology of clinical sodium glucose coâ€transport protein 2 inhibitors and relationship to suspected adverse drug reactions. Pharmacology Research and Perspectives, 2021, 9, e00867.	1.1	5
39	Sacubitril/valsartan in patients with symptomatic chronic heart failure with reduced ejection fraction. Journal of Prescribing Practice, 2019, 1, 182-192.	0.1	4
40	Voltammetric Behaviour of Drug Molecules as a Predictor of Metabolic Liabilities. Scientia Pharmaceutica, 2020, 88, 46.	0.7	4
41	Angiotensin II receptor blockers (ARBs) and manufacturing contamination: A retrospective National Register Study into suspected associated adverse drug reactions. British Journal of Clinical Pharmacology, 2022, 88, 4812-4827.	1.1	4
42	3-({5-Bromo-4-[pyrrolidin-1-yl]pyrimidin-2-yl}amino)phenol. MolBank, 2015, 2015, M859.	0.2	2
43	1,8-bis(2-hydroxy-3,5-di-tert-butylbenzyl)-4,11-dibenzyl-1,4,8,11-tetraazacyclotetradecane. MolBank, 2017, 2017, M963.	0.2	2
44	Aminothiazolones as potent, selective and cell active inhibitors of the PIM kinase family. Bioorganic and Medicinal Chemistry, 2020, 28, 115724.	1.4	1
45	Discovery and Characterization of a Cryptic Secondary Binding Site in the Molecular Chaperone HSP70. Molecules, 2022, 27, 817.	1.7	1
46	Unravelling the Interaction of Piperlongumine with the Nucleotide-Binding Domain of HSP70: A Spectroscopic and In Silico Study. Pharmaceuticals, 2021, 14, 1298.	1.7	1
47	A Sulfuryl Group Transfer Strategy to Selectively Prepare Sulfated Steroids and Isotopically Labelled Derivatives. Frontiers in Molecular Biosciences, 2021, 8, 776900.	1.6	1
48	43â€A novel role for small molecule glycomimetics in the protection against lipid-induced endothelial dysfunction. Heart, 2015, 101, A14.2-A14.	1.2	0
49	Investigations into the construction of the pentasubstituted ringCof Neosurugatoxin – a crystallographic study. Acta Crystallographica Section E: Crystallographic Communications, 2016, 72, 44-48.	0.2	O