

# Martin Scobie

## List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

28

papers

827

citations

14

h-index

28

g-index

34

ext. papers

1,085

ext. citations

9.9

avg, IF

3.01

L-index

#	Paper	IF	Citations
28	MTH1 inhibition eradicates cancer by preventing sanitation of the dNTP pool. <i>Nature</i> , <b>2014</b> , 508, 215-21	50.4	326
27	Cancer-Specific Synthetic Lethality between ATR and CHK1 Kinase Activities. <i>Cell Reports</i> , <b>2016</b> , 14, 298-308	30.8	72
26	Small-molecule inhibitor of OGG1 suppresses proinflammatory gene expression and inflammation. <i>Science</i> , <b>2018</b> , 362, 834-839	33.3	71
25	Rational design and validation of a Tip60 histone acetyltransferase inhibitor. <i>Scientific Reports</i> , <b>2014</b> , 4, 5372	4.9	70
24	Discovery of the First Potent and Selective Inhibitors of Human dCTP Pyrophosphatase 1. <i>Journal of Medicinal Chemistry</i> , <b>2016</b> , 59, 1140-1148	8.3	30
23	A new concise synthesis of 2,3-dihydroquinazolin-4(1H)-one derivatives. <i>New Journal of Chemistry</i> , <b>2013</b> , 37, 3595	3.6	30
22	A patient-derived xenograft pre-clinical trial reveals treatment responses and a resistance mechanism to karonudib in metastatic melanoma. <i>Cell Death and Disease</i> , <b>2018</b> , 9, 810	9.8	29
21	Targeted NUDT5 inhibitors block hormone signaling in breast cancer cells. <i>Nature Communications</i> , <b>2018</b> , 9, 250	17.4	28
20	Fragment-Based Discovery and Optimization of Enzyme Inhibitors by Docking of Commercial Chemical Space. <i>Journal of Medicinal Chemistry</i> , <b>2017</b> , 60, 8160-8169	8.3	25
19	Parallel synthesis of unsymmetrically substituted tetraphenyl porphyrins on Wang resin. <i>Tetrahedron Letters</i> , <b>2003</b> , 44, 5083-5086	2	19
18	dUTPase inhibition augments replication defects of 5-Fluorouracil. <i>Oncotarget</i> , <b>2017</b> , 8, 23713-23726	3.3	18
17	Application of combinatorial techniques in the synthesis of unsymmetrically substituted 5,15-diphenylporphyrins. <i>Tetrahedron Letters</i> , <b>2000</b> , 41, 2753-2757	2	17
16	Piperazin-1-ylpyridazine Derivatives Are a Novel Class of Human dCTP Pyrophosphatase 1 Inhibitors. <i>Journal of Medicinal Chemistry</i> , <b>2017</b> , 60, 4279-4292	8.3	14
15	A Convenient Microwave-Assisted Propylphosphonic Anhydride (T3P) Mediated One-Pot Pyrazolone Synthesis. <i>European Journal of Organic Chemistry</i> , <b>2013</b> , 2013, 5879-5885	3.2	14
14	Identification of Triazolothiadiazoles as Potent Inhibitors of the dCTP Pyrophosphatase 1. <i>Journal of Medicinal Chemistry</i> , <b>2017</b> , 60, 2148-2154	8.3	12
13	Computational and Experimental Druggability Assessment of Human DNA Glycosylases. <i>ACS Omega</i> , <b>2019</b> , 4, 11642-11656	3.9	11
12	A facile and efficient synthesis of tetrahydro- $\beta$ -carbolines. <i>Tetrahedron Letters</i> , <b>2013</b> , 54, 3554-3557	2	11

11	Targeting OGG1 arrests cancer cell proliferation by inducing replication stress. <i>Nucleic Acids Research</i> , <b>2020</b> , 48, 12234-12251	20.1	8
10	Development of a chemical probe against NUDT15. <i>Nature Chemical Biology</i> , <b>2020</b> , 16, 1120-1128	11.7	5
9	Diverse heterocyclic scaffolds as dCTP pyrophosphatase 1 inhibitors. Part 2: Pyridone- and pyrimidinone-derived systems. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2017</b> , 27, 3219-3225	2.9	3
8	Diverse heterocyclic scaffolds as dCTP pyrophosphatase 1 inhibitors. Part 1: Triazoles, triazolopyrimidines, triazinoindoles, quinoline hydrazones and arylpiperazines. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2017</b> , 27, 3897-3904	2.9	2
7	Rapid and Convenient Microwave-Assisted Synthesis of Primary Amines via Reductive N-Alkylation of Methyl Carbamate with Aldehydes. <i>Synthesis</i> , <b>2008</b> , 2008, 1679-1681	2.9	2
6	Crystal structures of NUDT15 variants enabled by a potent inhibitor reveal the structural basis for thiopurine sensitivity. <i>Journal of Biological Chemistry</i> , <b>2021</b> , 296, 100568	5.4	2
5	Pharmacological targeting of MTHFD2 suppresses acute myeloid leukemia by inducing thymidine depletion and replication stress.. <i>Nature Cancer</i> , <b>2022</b> , 3, 156-172	15.4	2
4	NUDT15-mediated hydrolysis limits the efficacy of anti-HCMV drug ganciclovir. <i>Cell Chemical Biology</i> , <b>2021</b> ,	8.2	1
3	MTH1 Inhibitors for the Treatment of Psoriasis. <i>Journal of Investigative Dermatology</i> , <b>2021</b> , 141, 2037-2048, e4	14.9	1
2	Structure-metabolism-relationships in the microsomal clearance of piperazin-1-ylpyridazines. <i>MedChemComm</i> , <b>2017</b> , 8, 1553-1560	5	0
1	MTH1 Inhibitor TH1579 Induces Oxidative DNA Damage and Mitotic Arrest in Acute Myeloid Leukemia. <i>Cancer Research</i> , <b>2021</b> , 81, 5733-5744	10.1	0