## Stephen Brand

List of Publications by Year in descending order

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516215 794141 19 960 16 19 citations g-index h-index papers 20 20 20 1184 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Reaction hijacking of tyrosine tRNA synthetase as a new whole-of-life-cycle antimalarial strategy. Science, 2022, 376, 1074-1079.	6.0	25
2	Discovery of Potent and Fast-Acting Antimalarial Bis-1,2,4-triazines. Journal of Medicinal Chemistry, 2021, 64, 4150-4162.	2.9	14
3	Scaffold-Hopping Strategy on a Series of Proteasome Inhibitors Led to a Preclinical Candidate for the Treatment of Visceral Leishmaniasis. Journal of Medicinal Chemistry, 2021, 64, 5905-5930.	2.9	25
4	Design of proteasome inhibitors with oral efficacy in vivo against <i>Plasmodium falciparum</i> and selectivity over the human proteasome. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	3.3	19
5	Optimisation of 2-(N-phenyl carboxamide) triazolopyrimidine antimalarials with moderate to slow acting erythrocytic stage activity. Bioorganic Chemistry, 2021, 115, 105244.	2.0	11
6	The proteasome as a target for protozoan parasites. Expert Opinion on Therapeutic Targets, 2019, 23, 903-914.	1.5	32
7	Preclinical candidate for the treatment of visceral leishmaniasis that acts through proteasome inhibition. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 9318-9323.	3.3	119
8	Identification of GSK3186899/DDD853651 as a Preclinical Development Candidate for the Treatment of Visceral Leishmaniasis. Journal of Medicinal Chemistry, 2019, 62, 1180-1202.	2.9	33
9	Pharmacological Validation of <i>N</i> -Myristoyltransferase as a Drug Target in <i>Leishmania donovani</i> . ACS Infectious Diseases, 2019, 5, 111-122.	1.8	55
10	A Molecular Hybridization Approach for the Design of Potent, Highly Selective, and Brain-Penetrant <i>N</i> -Myristoyltransferase Inhibitors. Journal of Medicinal Chemistry, 2018, 61, 8374-8389.	2.9	41
11	Chemical Validation of Methionyl-tRNA Synthetase as a Druggable Target in <i>Leishmania donovani</i> . ACS Infectious Diseases, 2017, 3, 718-727.	1.8	22
12	Discovery and Optimization of 5-Amino-1,2,3-triazole-4-carboxamide Series against <i>Trypanosoma cruzi</i> . Journal of Medicinal Chemistry, 2017, 60, 7284-7299.	2.9	31
13	Design and Synthesis of Brain Penetrant Trypanocidal <i>N</i> Journal of Medicinal Chemistry, 2017, 60, 9790-9806.	2.9	14
14	Validation of N-myristoyltransferase as Potential Chemotherapeutic Target in Mammal-Dwelling Stages of Trypanosoma cruzi. PLoS Neglected Tropical Diseases, 2016, 10, e0004540.	1.3	25
15	Development of Smallâ€Molecule <i>Trypanosoma brucei N</i> â€Myristoyltransferase Inhibitors: Discovery and Optimisation of a Novel Binding Mode. ChemMedChem, 2015, 10, 1821-1836.	1.6	20
16	Development of a Fluorescence-based Trypanosoma cruzi CYP51 Inhibition Assay for Effective Compound Triaging in Drug Discovery Programmes for Chagas Disease. PLoS Neglected Tropical Diseases, 2015, 9, e0004014.	1.3	43
17	Lead Optimization of a Pyrazole Sulfonamide Series of <i>Trypanosoma bruceiN</i> -Myristoyltransferase Inhibitors: Identification and Evaluation of CNS Penetrant Compounds as Potential Treatments for Stage 2 Human African Trypanosomiasis. Journal of Medicinal Chemistry, 2014, 57, 9855-9869.	2.9	57
18	Discovery of a Novel Class of Orally Active Trypanocidal <i>N</i> Journal of Medicinal Chemistry, 2012, 55, 140-152.	2.9	102

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#	Article	lF	CITATIONS
19	N-myristoyltransferase inhibitors as new leads to treat sleeping sickness. Nature, 2010, 464, 728-732.	13.7	272