Conor R Caffrey

List of Publications by Year in descending order

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		159585	155660
86	3,564	30	55
papers	citations	h-index	g-index
124	124	124	3566
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Brazilian green propolis reduces worm burden and hepatic granuloma formation in a Schistosoma mansoni experimental murine model. Parasitology Research, 2022, 121, 775-780.	1.6	2
2	Biomechanical interactions of Schistosoma mansoni eggs with vascular endothelial cells facilitate egg extravasation. PLoS Pathogens, 2022, 18, e1010309.	4.7	3
3	Druggable Hot Spots in the Schistosomiasis Cathepsin B1 Target Identified by Functional and Binding Mode Analysis of Potent Vinyl Sulfone Inhibitors. ACS Infectious Diseases, 2021, 7, 1077-1088.	3.8	9
4	Azanitrile Inhibitors of the SmCB1 Protease Target Are Lethal to <i>Schistosoma mansoni</i> : Structural and Mechanistic Insights into Chemotype Reactivity. ACS Infectious Diseases, 2021, 7, 189-201.	3.8	9
5	Congeners Derived from Microtubule-Active Phenylpyrimidines Produce a Potent and Long-Lasting Paralysis of <i>Schistosoma mansoni</i> In Vitro. ACS Infectious Diseases, 2021, 7, 1089-1103.	3.8	6
6	Understanding the key processes of excellence as a prerequisite to establishing academic centres of excellence in Africa. BMC Medical Education, 2021, 21, 36.	2.4	7
7	A Machine Learning Strategy for Drug Discovery Identifies Anti-Schistosomal Small Molecules. ACS Infectious Diseases, 2021, 7, 406-420.	3.8	18
8	Lead Optimization of 3,5-Disubstituted-7-Azaindoles for the Treatment of Human African Trypanosomiasis. Journal of Medicinal Chemistry, 2021, 64, 9404-9430.	6.4	6
9	Antiparasitic Properties of Propolis Extracts and Their Compounds. Chemistry and Biodiversity, 2021, 18, e2100310.	2.1	13
10	Structure-Based Optimization of Quinazolines as Cruzain and <i>Tbr</i> CATL Inhibitors. Journal of Medicinal Chemistry, 2021, 64, 13054-13071.	6.4	19
11	Should the enzyme name â€~rhodesain' be discontinued?. Molecular and Biochemical Parasitology, 2021, 245, 111395.	1.1	8
12	Anti-schistosomal activities of quinoxaline-containing compounds: From hit identification to lead optimisation. European Journal of Medicinal Chemistry, 2021, 226, 113823.	5.5	8
13	Hit-to-Lead Optimization of Benzoxazepinoindazoles As Human African Trypanosomiasis Therapeutics. Journal of Medicinal Chemistry, 2020, 63, 2527-2546.	6.4	11
14	Selectivity and Physicochemical Optimization of Repurposed Pyrazolo[1,5- <i>b</i>]pyridazines for the Treatment of Human African Trypanosomiasis. Journal of Medicinal Chemistry, 2020, 63, 756-783.	6.4	10
15	A single-cell RNA-seq atlas of <i>Schistosoma mansoni</i> identifies a key regulator of blood feeding. Science, 2020, 369, 1644-1649.	12.6	108
16	Uncovering Biological Application of Brazilian Green Propolis: A Phenotypic Screening against Schistosoma mansoni. Chemistry and Biodiversity, 2020, 17, e2000277.	2.1	3
17	Identification of anisomycin, prodigiosin and obatoclax as compounds with broad-spectrum anti-parasitic activity. PLoS Neglected Tropical Diseases, 2020, 14, e0008150.	3.0	20
18	Isoforms of Cathepsin B1 in Neurotropic Schistosomula of Trichobilharzia regenti Differ in Substrate Preferences and a Highly Expressed Catalytically Inactive Paralog Binds Cystatin. Frontiers in Cellular and Infection Microbiology, 2020, 10, 66.	3.9	3

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19	Design, synthesis, and <i>inÂvitro</i> evaluation of aza-peptide aldehydes and ketones as novel and selective protease inhibitors. Journal of Enzyme Inhibition and Medicinal Chemistry, 2020, 35, 1387-1402.	5.2	6
20	Efficacy, metabolism and pharmacokinetics of Ro 15-5458, a forgotten schistosomicidal 9-acridanone hydrazone. Journal of Antimicrobial Chemotherapy, 2020, 75, 2925-2932.	3.0	3
21	Synthesis and Bioactivity of Phthalimide Analogs as Potential Drugs to Treat Schistosomiasis, a Neglected Disease of Poverty. Pharmaceuticals, 2020, 13, 25.	3.8	9
22	Novel and selective inactivators of Triosephosphate isomerase with anti-trematode activity. Scientific Reports, 2020, 10, 2587.	3.3	12
23	Structure–Bioactivity Relationships of Lapatinib Derived Analogs against <i>Schistosoma mansoni</i> . ACS Medicinal Chemistry Letters, 2020, 11, 258-265.	2.8	2
24	A multi-dimensional, time-lapse, high content screening platform applied to schistosomiasis drug discovery. Communications Biology, 2020, 3, 747.	4.4	16
25	Title is missing!. , 2020, 14, e0008150.		0
26	Title is missing!. , 2020, 14, e0008150.		0
27	Title is missing!. , 2020, 14, e0008150.		0
28	Title is missing!. , 2020, 14, e0008150.		0
29	Development and optimization of a high-throughput screening method utilizing Ancylostoma ceylanicum egg hatching to identify novel anthelmintics. PLoS ONE, 2019, 14, e0217019.	2.5	16
30	The Proteasome as a Drug Target in the Metazoan Pathogen, <i>Schistosoma mansoni</i> . ACS Infectious Diseases, 2019, 5, 1802-1812.	3.8	25
31	Molecular characterization and functional analysis of the Schistosoma mekongi Ca2+-dependent cysteine protease (calpain). Parasites and Vectors, 2019, 12, 383.	2.5	13
32	Benzimidazole inhibitors of the major cysteine protease of <i>Trypanosoma brucei</i> . Future Medicinal Chemistry, 2019, 11, 1537-1551.	2.3	7
33	A secreted schistosome cathepsin B1 cysteine protease and acute schistosome infection induce a transient T helper 17 response. PLoS Neglected Tropical Diseases, 2019, 13, e0007070.	3.0	20
34	Discovery and characterization of trypanocidal cysteine protease inhibitors from the â€~malaria box'. European Journal of Medicinal Chemistry, 2019, 179, 765-778.	5.5	19
35	Bioactivity of Farnesyltransferase Inhibitors Against Entamoeba histolytica and Schistosoma mansoni. Frontiers in Cellular and Infection Microbiology, 2019, 9, 180.	3.9	12
36	Evaluation of a class of isatinoids identified from a high-throughput screen of human kinase inhibitors as anti-Sleeping Sickness agents. PLoS Neglected Tropical Diseases, 2019, 13, e0007129.	3.0	4

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37	Quantifying the mechanics of locomotion of the schistosome pathogen with respect to changes in its physical environment. Journal of the Royal Society Interface, 2019, 16, 20180675.	3.4	13
38	Multi-center screening of the Pathogen Box collection for schistosomiasis drug discovery. Parasites and Vectors, 2019, 12, 493.	2.5	20
39	High Throughput and Computational Repurposing for Neglected Diseases. Pharmaceutical Research, 2019, 36, 27.	3.5	37
40	Sertraline, Paroxetine, and Chlorpromazine Are Rapidly Acting Anthelmintic Drugs Capable of Clinical Repurposing. Scientific Reports, 2018, 8, 975.	3.3	64
41	Octopamine-signaling in the metazoan pathogen, Schistosoma mansoni: localization, small-molecule screening and opportunities for drug development. DMM Disease Models and Mechanisms, 2018, 11, .	2.4	6
42	TPT sulfonate, a single, oral dose schistosomicidal prodrug: In vivo efficacy, disposition and metabolic profiling. International Journal for Parasitology: Drugs and Drug Resistance, 2018, 8, 571-586.	3.4	13
43	Cysteine proteases during larval migration and development of helminths in their final host. PLoS Neglected Tropical Diseases, 2018, 12, e0005919.	3.0	27
44	Brainâ€Penetrant Triazolopyrimidine and Phenylpyrimidine Microtubule Stabilizers as Potential Leads to Treat Human African Trypanosomiasis. ChemMedChem, 2018, 13, 1751-1754.	3.2	19
45	SmSP2: A serine protease secreted by the blood fluke pathogen Schistosoma mansoni with anti-hemostatic properties. PLoS Neglected Tropical Diseases, 2018, 12, e0006446.	3.0	26
46	Substrate Specificity of Cysteine Proteases Beyond the S2 Pocket: Mutagenesis and Molecular Dynamics Investigation of Fasciola hepatica Cathepsins L. Frontiers in Molecular Biosciences, 2018, 5, 40.	3.5	10
47	Effect of Phenotypic Screening of Extracts and Fractions of <i> Erythrophleum ivorense</i> Leaf and Stem Bark on Immature and Adult Stages of <i> Schistosoma mansoni</i> . Journal of Parasitology Research, 2018, 2018, 1-7.	1.2	13
48	Cysteine proteases as digestive enzymes in parasitic helminths. PLoS Neglected Tropical Diseases, 2018, 12, e0005840.	3.0	82
49	Targeting proteasomes in infectious organisms to combat disease. FEBS Journal, 2017, 284, 1503-1517.	4.7	40
50	Phenotypic, chemical and functional characterization of cyclic nucleotide phosphodiesterase 4 (PDE4) as a potential anthelmintic drug target. PLoS Neglected Tropical Diseases, 2017, 11, e0005680.	3.0	36
51	Odanacatib, a Cathepsin K Cysteine Protease Inhibitor, Kills Hookworm In Vivo. Pharmaceuticals, 2016, 9, 39.	3.8	7
52	Open Source Drug Discovery with the Malaria Box Compound Collection for Neglected Diseases and Beyond. PLoS Pathogens, 2016, 12, e1005763.	4.7	244
53	Structure-Bioactivity Relationship for Benzimidazole Thiophene Inhibitors of Polo-Like Kinase 1 (PLK1), a Potential Drug Target in Schistosoma mansoni. PLoS Neglected Tropical Diseases, 2016, 10, e0004356.	3.0	56
54	Evaluation of the CCA Immuno-Chromatographic Test to Diagnose Schistosoma mansoni in Minas Gerais State, Brazil. PLoS Neglected Tropical Diseases, 2016, 10, e0004357.	3.0	18

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55	Sex-Biased Transcriptome of Schistosoma mansoni: Host-Parasite Interaction, Genetic Determinants and Epigenetic Regulators Are Associated with Sexual Differentiation. PLoS Neglected Tropical Diseases, 2016, 10, e0004930.	3.0	57
56	Excretion/secretion products from Schistosoma mansoni adults, eggs and schistosomula have unique peptidase specificity profiles. Biochimie, 2016, 122, 99-109.	2.6	31
57	Prolyl Oligopeptidase from the Blood Fluke Schistosoma mansoni: From Functional Analysis to Anti-schistosomal Inhibitors. PLoS Neglected Tropical Diseases, 2015, 9, e0003827.	3.0	34
58	The QDREC web server: determining dose–response characteristics of complex macroparasites in phenotypic drug screens. Bioinformatics, 2015, 31, 1515-1518.	4.1	21
59	Synthesis of a Sugar-Based Thiosemicarbazone Series and Structure-Activity Relationship versus the Parasite Cysteine Proteases Rhodesain, Cruzain, and Schistosoma mansoni Cathepsin B1. Antimicrobial Agents and Chemotherapy, 2015, 59, 2666-2677.	3.2	57
60	Caenorhabditis elegans is a useful model for anthelmintic discovery. Nature Communications, 2015, 6, 7485.	12.8	163
61	Regulation of Schistosoma mansoni Development and Reproduction by the Mitogen-Activated Protein Kinase Signaling Pathway. PLoS Neglected Tropical Diseases, 2014, 8, e2949.	3.0	73
62	Trypsin- and Chymotrypsin-Like Serine Proteases in Schistosoma mansoni – â€~The Undiscovered Country'. PLoS Neglected Tropical Diseases, 2014, 8, e2766.	3.0	31
63	Serum albumin and α-1 acid glycoprotein impede the killing of Schistosoma mansoni by the tyrosine kinase inhibitor Imatinib. International Journal for Parasitology: Drugs and Drug Resistance, 2014, 4, 287-295.	3.4	34
64	Activation Route of the Schistosoma mansoni Cathepsin B1 Drug Target: Structural Map with a Glycosaminoglycan Switch. Structure, 2014, 22, 1786-1798.	3.3	34
65	Chemical and Genetic Validation of the Statin Drug Target to Treat the Helminth Disease, Schistosomiasis. PLoS ONE, 2014, 9, e87594.	2.5	62
66	Cure of Hookworm Infection with a Cysteine Protease Inhibitor. PLoS Neglected Tropical Diseases, 2012, 6, e1680.	3.0	28
67	Mapping the Pro-Peptide of the <i>Schistosoma mansoni</i> Cathepsin B1 Drug Target: Modulation of Inhibition by Heparin and Design of Mimetic Inhibitors. ACS Chemical Biology, 2011, 6, 609-617.	3.4	34
68	Structural Basis for Inhibition of Cathepsin B Drug Target from the Human Blood Fluke, Schistosoma mansoni. Journal of Biological Chemistry, 2011, 286, 35770-35781.	3.4	60
69	RNA Interference in Schistosoma mansoni Schistosomula: Selectivity, Sensitivity and Operation for Larger-Scale Screening. PLoS Neglected Tropical Diseases, 2010, 4, e850.	3.0	107
70	Chapter 4 Peptidases of Trematodes. Advances in Parasitology, 2009, 69, 205-297.	3.2	70
71	SmCL3, a Gastrodermal Cysteine Protease of the Human Blood Fluke Schistosoma mansoni. PLoS Neglected Tropical Diseases, 2009, 3, e449.	3.0	45
72	Differential use of protease families for invasion by schistosome cercariae. Biochimie, 2008, 90, 345-358.	2.6	100

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73	Schistosomiasis Mansoni: Novel Chemotherapy Using a Cysteine Protease Inhibitor. PLoS Medicine, 2007, 4, e14.	8.4	229
74	Chemical Composition and Cruzain Inhibitory Activity of <i>Croton draco</i> Bark Essential Oil from Monteverde, Costa Rica. Natural Product Communications, 2007, 2, 1934578X0700200.	0.5	4
75	Cruzain Inhibitory Activity of Leaf Essential Oils of Neotropical Lauraceae and Essential Oil Components. Natural Product Communications, 2007, 2, 1934578X0700201.	0.5	18
76	3-O-(3′-Hydroxytetradecanoyl)lupeol from Sorocea trophoides Inhibits Cruzain. Natural Product Communications, 2007, 2, 1934578X0700200.	0.5	6
77	Inhibition of Cruzain by Triterpenoids Isolated from a Salacia Species from Monteverde, Costa Rica. Natural Product Communications, 2007, 2, 1934578X0700201.	0.5	5
78	Chemotherapy of schistosomiasis: present and future. Current Opinion in Chemical Biology, 2007, 11, 433-439.	6.1	251
79	A Multienzyme Network Functions in Intestinal Protein Digestion by a Platyhelminth Parasite. Journal of Biological Chemistry, 2006, 281, 39316-39329.	3.4	214
80	Multiple cathepsin B isoforms in schistosomula of Trichobilharzia regenti: identification, characterisation and putative role in migration and nutrition. International Journal for Parasitology, 2005, 35, 895-910.	3.1	50
81	Blood â€~n' guts: an update on schistosome digestive peptidases. Trends in Parasitology, 2004, 20, 241-248.	3.3	147
82	Functional expression and characterization of Schistosoma mansoni cathepsin B and its trans-activation by an endogenous asparaginyl endopeptidase. Molecular and Biochemical Parasitology, 2003, 131, 65-75.	1.1	147
83	Screening of acyl hydrazide proteinase inhibitors for antiparasitic activity against Trypanosoma brucei. International Journal of Antimicrobial Agents, 2002, 19, 227-231.	2.5	30
84	SmCB2, a novel tegumental cathepsin B from adult Schistosoma mansoni. Molecular and Biochemical Parasitology, 2002, 121, 49-61.	1.1	69
85	Identification of a cDNA encoding an active asparaginyl endopeptidase ofSchistosoma mansoniand its expression inPichia pastoris1. FEBS Letters, 2000, 466, 244-248.	2.8	64
86	Discovery of pH-Selective Marine and Plant Natural Product Inhibitors of Cathepsin B Revealed by Screening at Acidic and Neutral pH Conditions. ACS Omega, 0, , .	3.5	2