

Conor R Caffrey

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118
papers

2,604
citations

27
h-index

49
g-index

124
ext. papers

3,189
ext. citations

5.2
avg, IF

4.92
L-index

#	Paper	IF	Citations
118	Chemotherapy of schistosomiasis: present and future. <i>Current Opinion in Chemical Biology</i> , 2007 , 11, 433-9	9.7	226
117	Schistosomiasis mansoni: novel chemotherapy using a cysteine protease inhibitor. <i>PLoS Medicine</i> , 2007 , 4, e14	11.6	199
116	A multienzyme network functions in intestinal protein digestion by a platyhelminth parasite. <i>Journal of Biological Chemistry</i> , 2006 , 281, 39316-29	5.4	180
115	Open Source Drug Discovery with the Malaria Box Compound Collection for Neglected Diseases and Beyond. <i>PLoS Pathogens</i> , 2016 , 12, e1005763	7.6	167
114	Functional expression and characterization of <i>Schistosoma mansoni</i> cathepsin B and its trans-activation by an endogenous asparaginyl endopeptidase. <i>Molecular and Biochemical Parasitology</i> , 2003 , 131, 65-75	1.9	128
113	Blood Tguts: an update on schistosome digestive peptidases. <i>Trends in Parasitology</i> , 2004 , 20, 241-8	6.4	125
112	<i>Caenorhabditis elegans</i> is a useful model for anthelmintic discovery. <i>Nature Communications</i> , 2015 , 6, 7485	17.4	103
111	RNA interference in <i>Schistosoma mansoni</i> schistosomula: selectivity, sensitivity and operation for larger-scale screening. <i>PLoS Neglected Tropical Diseases</i> , 2010 , 4, e850	4.8	95
110	Differential use of protease families for invasion by schistosome cercariae. <i>Biochimie</i> , 2008 , 90, 345-58	4.6	87
109	SmCB2, a novel tegumental cathepsin B from adult <i>Schistosoma mansoni</i> . <i>Molecular and Biochemical Parasitology</i> , 2002 , 121, 49-61	1.9	63
108	Chapter 4. Peptidases of trematodes. <i>Advances in Parasitology</i> , 2009 , 69, 205-97	3.2	59
107	Identification of a cDNA encoding an active asparaginyl endopeptidase of <i>Schistosoma mansoni</i> and its expression in <i>Pichia pastoris</i> . <i>FEBS Letters</i> , 2000 , 466, 244-8	3.8	52
106	Structural basis for inhibition of cathepsin B drug target from the human blood fluke, <i>Schistosoma mansoni</i> . <i>Journal of Biological Chemistry</i> , 2011 , 286, 35770-35781	5.4	51
105	Chemical and genetic validation of the statin drug target to treat the helminth disease, schistosomiasis. <i>PLoS ONE</i> , 2014 , 9, e87594	3.7	51
104	Synthesis of a sugar-based thiosemicarbazone series and structure-activity relationship versus the parasite cysteine proteases rhodesain, cruzain, and <i>Schistosoma mansoni</i> cathepsin B1. <i>Antimicrobial Agents and Chemotherapy</i> , 2015 , 59, 2666-77	5.9	45
103	Cysteine proteases as digestive enzymes in parasitic helminths. <i>PLoS Neglected Tropical Diseases</i> , 2018 , 12, e0005840	4.8	43
102	SmCL3, a gastrodermal cysteine protease of the human blood fluke <i>Schistosoma mansoni</i> . <i>PLoS Neglected Tropical Diseases</i> , 2009 , 3, e449	4.8	42

101	Regulation of <i>Schistosoma mansoni</i> development and reproduction by the mitogen-activated protein kinase signaling pathway. <i>PLoS Neglected Tropical Diseases</i> , 2014 , 8, e2949	4.8	40
100	Multiple cathepsin B isoforms in schistosomula of <i>Trichobilharzia regenti</i> : identification, characterisation and putative role in migration and nutrition. <i>International Journal for Parasitology</i> , 2005 , 35, 895-910	4.3	40
99	Structure-Bioactivity Relationship for Benzimidazole Thiophene Inhibitors of Polo-Like Kinase 1 (PLK1), a Potential Drug Target in <i>Schistosoma mansoni</i> . <i>PLoS Neglected Tropical Diseases</i> , 2016 , 10, e0004356	4.8	40
98	Sertraline, Paroxetine, and Chlorpromazine Are Rapidly Acting Anthelmintic Drugs Capable of Clinical Repurposing. <i>Scientific Reports</i> , 2018 , 8, 975	4.9	39
97	Sex-Biased Transcriptome of <i>Schistosoma mansoni</i> : Host-Parasite Interaction, Genetic Determinants and Epigenetic Regulators Are Associated with Sexual Differentiation. <i>PLoS Neglected Tropical Diseases</i> , 2016 , 10, e0004930	4.8	38
96	A single-cell RNA-seq atlas of identifies a key regulator of blood feeding. <i>Science</i> , 2020 , 369, 1644-1649	33.3	35
95	High Throughput and Computational Repurposing for Neglected Diseases. <i>Pharmaceutical Research</i> , 2018 , 36, 27	4.5	33
94	Targeting proteasomes in infectious organisms to combat disease. <i>FEBS Journal</i> , 2017 , 284, 1503-1517	5.7	32
93	Phenotypic, chemical and functional characterization of cyclic nucleotide phosphodiesterase 4 (PDE4) as a potential anthelmintic drug target. <i>PLoS Neglected Tropical Diseases</i> , 2017 , 11, e0005680	4.8	29
92	Serum albumin and β_2 acid glycoprotein impede the killing of <i>Schistosoma mansoni</i> by the tyrosine kinase inhibitor Imatinib. <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2014 , 4, 287-95	4	29
91	Prolyl Oligopeptidase from the Blood Fluke <i>Schistosoma mansoni</i> : From Functional Analysis to Anti-schistosomal Inhibitors. <i>PLoS Neglected Tropical Diseases</i> , 2015 , 9, e0003827	4.8	26
90	Screening of acyl hydrazide proteinase inhibitors for antiparasitic activity against <i>Trypanosoma brucei</i> . <i>International Journal of Antimicrobial Agents</i> , 2002 , 19, 227-31	14.3	25
89	Trypsin- and Chymotrypsin-like serine proteases in <i>Schistosoma mansoni</i> -- The undiscovered country? <i>PLoS Neglected Tropical Diseases</i> , 2014 , 8, e2766	4.8	24
88	Mapping the pro-peptide of the <i>Schistosoma mansoni</i> cathepsin B1 drug target: modulation of inhibition by heparin and design of mimetic inhibitors. <i>ACS Chemical Biology</i> , 2011 , 6, 609-17	4.9	24
87	Cure of hookworm infection with a cysteine protease inhibitor. <i>PLoS Neglected Tropical Diseases</i> , 2012 , 6, e1680	4.8	24
86	Excretion/secretion products from <i>Schistosoma mansoni</i> adults, eggs and schistosomula have unique peptidase specificity profiles. <i>Biochimie</i> , 2016 , 122, 99-109	4.6	23
85	Activation route of the <i>Schistosoma mansoni</i> cathepsin B1 drug target: structural map with a glycosaminoglycan switch. <i>Structure</i> , 2014 , 22, 1786-1798	5.2	23
84	Cysteine proteases during larval migration and development of helminths in their final host. <i>PLoS Neglected Tropical Diseases</i> , 2018 , 12, e0005919	4.8	18

83	SmSP2: A serine protease secreted by the blood fluke pathogen <i>Schistosoma mansoni</i> with anti-hemostatic properties. <i>PLoS Neglected Tropical Diseases</i> , 2018 , 12, e0006446	4.8	17
82	Evaluation of the CCA Immuno-Chromatographic Test to Diagnose <i>Schistosoma mansoni</i> in Minas Gerais State, Brazil. <i>PLoS Neglected Tropical Diseases</i> , 2016 , 10, e0004357	4.8	16
81	The Proteasome as a Drug Target in the Metazoan Pathogen,. <i>ACS Infectious Diseases</i> , 2019 , 5, 1802-1813	3.5	15
80	The QDREC web server: determining dose-response characteristics of complex macroparasites in phenotypic drug screens. <i>Bioinformatics</i> , 2015 , 31, 1515-8	7.2	14
79	Cruzain Inhibitory Activity of Leaf Essential Oils of Neotropical Lauraceae and Essential Oil Components. <i>Natural Product Communications</i> , 2007 , 2, 1934578X0700201	0.9	12
78	Brain-Penetrant Triazolopyrimidine and Phenylpyrimidine Microtubule Stabilizers as Potential Leads to Treat Human African Trypanosomiasis. <i>ChemMedChem</i> , 2018 , 13, 1751-1754	3.7	11
77	Multi-center screening of the Pathogen Box collection for schistosomiasis drug discovery. <i>Parasites and Vectors</i> , 2019 , 12, 493	4	10
76	A secreted schistosome cathepsin B1 cysteine protease and acute schistosome infection induce a transient T helper 17 response. <i>PLoS Neglected Tropical Diseases</i> , 2019 , 13, e0007070	4.8	9
75	Effect of Phenotypic Screening of Extracts and Fractions of Leaf and Stem Bark on Immature and Adult Stages of. <i>Journal of Parasitology Research</i> , 2018 , 2018, 9431467	1.9	9
74	Drug Discovery and Development for Schistosomiasis. <i>Methods and Principles in Medicinal Chemistry</i> , 2019 , 187-225	0.4	8
73	Identification of anisomycin, prodigiosin and obatoclax as compounds with broad-spectrum anti-parasitic activity. <i>PLoS Neglected Tropical Diseases</i> , 2020 , 14, e0008150	4.8	8
72	A single-cell RNAseq atlas of the pathogenic stage of <i>Schistosoma mansoni</i> identifies a key regulator of blood feeding		8
71	A Machine Learning Strategy for Drug Discovery Identifies Anti-Schistosomal Small Molecules. <i>ACS Infectious Diseases</i> , 2021 , 7, 406-420	5.5	8
70	Discovery and characterization of trypanocidal cysteine protease inhibitors from the <i>T. malariae</i> boxT. <i>European Journal of Medicinal Chemistry</i> , 2019 , 179, 765-778	6.8	7
69	Integrating and Mining Helminth Genomes to Discover and Prioritize Novel Therapeutic Targets. 2012 , 43-59		7
68	Bioactivity of Farnesyltransferase Inhibitors Against and. <i>Frontiers in Cellular and Infection Microbiology</i> , 2019 , 9, 180	5.9	6
67	Quantifying the mechanics of locomotion of the schistosome pathogen with respect to changes in its physical environment. <i>Journal of the Royal Society Interface</i> , 2019 , 16, 20180675	4.1	6
66	Substrate Specificity of Cysteine Proteases Beyond the S Pocket: Mutagenesis and Molecular Dynamics Investigation of Cathepsins L. <i>Frontiers in Molecular Biosciences</i> , 2018 , 5, 40	5.6	6

65	Antinematodal Drugs [Modes of Action and Resistance: And Worms Will Not Come to Thee (Shakespeare: Cymbeline: IV, ii) 2012 , 233-249		6
64	Hit-to-Lead Optimization of Benzoxazepinoindazoles As Human African Trypanosomiasis Therapeutics. <i>Journal of Medicinal Chemistry</i> , 2020 , 63, 2527-2546	8.3	6
63	Selectivity and Physicochemical Optimization of Repurposed Pyrazolo[1,5-]pyridazines for the Treatment of Human African Trypanosomiasis. <i>Journal of Medicinal Chemistry</i> , 2020 , 63, 756-783	8.3	6
62	Odanacatib, a Cathepsin K Cysteine Protease Inhibitor, Kills Hookworm In Vivo. <i>Pharmaceuticals</i> , 2016 , 9,	5.2	6
61	TPT sulfonate, a single, oral dose schistosomicidal prodrug: In vivo efficacy, disposition and metabolic profiling. <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2018 , 8, 571-586	4	6
60	Structure-Based Optimization of Quinazolines as Cruzain and CATL Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 13054-13071	8.3	6
59	Use of Rodent Models in the Discovery of Novel Anthelmintics 181-199		6
58	Benzimidazole inhibitors of the major cysteine protease of. <i>Future Medicinal Chemistry</i> , 2019 , 11, 1537-1551	4.5	5
57	Novel and selective inactivators of Triosephosphate isomerase with anti-trematode activity. <i>Scientific Reports</i> , 2020 , 10, 2587	4.9	5
56	Molecular characterization and functional analysis of the Schistosoma mekongi Ca-dependent cysteine protease (calpain). <i>Parasites and Vectors</i> , 2019 , 12, 383	4	5
55	Mechanism-Based Screening Strategies for Anthelmintic Discovery 2012 , 121-134		5
54	Discovery, Mode of Action, and Commercialization of Derquantel 2012 , 297-307		5
53	Prospects for Immunoprophylaxis Against Fasciola hepatica (Liver Fluke) 2012 , 465-484		5
52	Promise of Bacillus thuringiensis Crystal Proteins as Anthelmintics 2012 , 267-281		5
51	3-O-(3?-Hydroxytetradecanoyl)lupeol from Sorocea trophoides Inhibits Cruzain. <i>Natural Product Communications</i> , 2007 , 2, 1934578X0700200	0.9	5
50	A multi-dimensional, time-lapse, high content screening platform applied to schistosomiasis drug discovery. <i>Communications Biology</i> , 2020 , 3, 747	6.7	5
49	Octopamine signaling in the metazoan pathogen : localization, small-molecule screening and opportunities for drug development. <i>DMM Disease Models and Mechanisms</i> , 2018 , 11,	4.1	5
48	Synthesis and Bioactivity of Phthalimide Analogs as Potential Drugs to Treat Schistosomiasis, a Neglected Disease of Poverty. <i>Pharmaceuticals</i> , 2020 , 13,	5.2	4

47	Development and optimization of a high-throughput screening method utilizing <i>Ancylostoma ceylanicum</i> egg hatching to identify novel anthelmintics. <i>PLoS ONE</i> , 2019 , 14, e0217019	3.7	4
46	Drug Discovery Approaches toward Anti-Parasitic Agents 2011 , 1-20		4
45	How Relevant is <i>Caenorhabditis elegans</i> as a Model for the Analysis of Parasitic Nematode Biology? 2012 , 23-41		4
44	Inhibition of Cruzain by Triterpenoids Isolated from a <i>Salacia</i> Species from Monteverde, Costa Rica. <i>Natural Product Communications</i> , 2007 , 2, 1934578X0700201	0.9	4
43	Azanitrile Inhibitors of the SmCB1 Protease Target Are Lethal to : Structural and Mechanistic Insights into Chemotype Reactivity. <i>ACS Infectious Diseases</i> , 2021 , 7, 189-201	5.5	4
42	Evaluation of a class of isatinoids identified from a high-throughput screen of human kinase inhibitors as anti-Sleeping Sickness agents. <i>PLoS Neglected Tropical Diseases</i> , 2019 , 13, e0007129	4.8	3
41	Efficacy, metabolism and pharmacokinetics of Ro 15-5458, a forgotten schistosomicidal 9-acridanone hydrazone. <i>Journal of Antimicrobial Chemotherapy</i> , 2020 , 75, 2925-2932	5.1	3
40	Chemical Composition and Cruzain Inhibitory Activity of <i>Croton draco</i> Bark Essential Oil from Monteverde, Costa Rica. <i>Natural Product Communications</i> , 2007 , 2, 1934578X0700200	0.9	3
39	Congeners Derived from Microtubule-Active Phenylpyrimidines Produce a Potent and Long-Lasting Paralysis of In Vitro. <i>ACS Infectious Diseases</i> , 2021 , 7, 1089-1103	5.5	3
38	Antiparasitic Properties of Propolis Extracts and Their Compounds. <i>Chemistry and Biodiversity</i> , 2021 , 18, e2100310	2.5	3
37	Ligand-Gated Ion Channels as Targets for Anthelmintic Drugs: Past, Current, and Future Perspectives1-21		3
36	Anthelmintic Drugs: Tools and Shortcuts for the Long Road From Discovery to Product217-232		3
35	Isoforms of Cathepsin B1 in Neurotropic Schistosomula of Differ in Substrate Preferences and a Highly Expressed Catalytically Inactive Paralog Binds Cystatin. <i>Frontiers in Cellular and Infection Microbiology</i> , 2020 , 10, 66	5.9	2
34	Chemotherapeutic Development Strategies for Schistosomiasis299-321		2
33	Druggable Hot Spots in the Schistosomiasis Cathepsin B1 Target Identified by Functional and Binding Mode Analysis of Potent Vinyl Sulfone Inhibitors. <i>ACS Infectious Diseases</i> , 2021 , 7, 1077-1088	5.5	2
32	Understanding the key processes of excellence as a prerequisite to establishing academic centres of excellence in Africa. <i>BMC Medical Education</i> , 2021 , 21, 36	3.3	2
31	Quantitative High-Content Screening-Based Drug Discovery against Helminthic Diseases159-179		2
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27	Identification and Profiling of Nematicidal Compounds in Veterinary Parasitology135-157		2
26	Uncovering Biological Application of Brazilian Green Propolis: A Phenotypic Screening against <i>Schistosoma mansoni</i> . <i>Chemistry and Biodiversity</i> , 2020 , 17, e2000277	2.5	1
25	Design, synthesis, and evaluation of aza-peptide aldehydes and ketones as novel and selective protease inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1387-1402	5.6	1
24	Structure-Bioactivity Relationships of Lapatinib Derived Analogs against. <i>ACS Medicinal Chemistry Letters</i> , 2020 , 11, 258-265	4.3	1
23	Barefoot thruTthe Valley of Darkness: Preclinical Development of a Human Hookworm Vaccine341-356		1
22	Lead Optimization of 3,5-Disubstituted-7-Azaindoles for the Treatment of Human African Trypanosomiasis. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 9404-9430	8.3	1
21	Should the enzyme name ThodesainTbe discontinued?. <i>Molecular and Biochemical Parasitology</i> , 2021 , 245, 111395	1.9	1
20	Anti-schistosomal activities of quinoxaline-containing compounds: From hit identification to lead optimisation. <i>European Journal of Medicinal Chemistry</i> , 2021 , 226, 113823	6.8	1
19	Monepantel: From Discovery to Mode of Action283-296		1
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15	Biomechanical interactions of <i>Schistosoma mansoni</i> eggs with vascular endothelial cells facilitate egg extravasation.. <i>PLoS Pathogens</i> , 2022 , 18, e1010309	7.6	0
14	Anthelmintic drug discovery: Into the future 2016 , 215-228		
13	Mechanisms of Immune Modulation by <i>Fasciola hepatica</i> : Importance for Vaccine Development and for Novel Immunotherapeutics 2012 , 451-463		
12	RNA Interference as a Tool for Drug Discovery in Parasitic Flatworms 2012 , 105-119		

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- 3 Identification of anisomycin, prodigiosin and obatoclax as compounds with broad-spectrum anti-parasitic activity **2020**, 14, e0008150
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- 1 Identification of anisomycin, prodigiosin and obatoclax as compounds with broad-spectrum anti-parasitic activity **2020**, 14, e0008150