Pascal MÃ¤ser

List of Publications by Year
in descending order

118 papers	4,361 citations	126708 33 h-index	118652 62 g-index
122 all docs	122 docs citations	122 times ranked	4754 citing authors

#	Article	IF	Citations
1	Laboratory Selection of Trypanosomatid Pathogens for Drug Resistance. Pharmaceuticals, 2022, 15, 135.	1.7	1
2	Multi-insecticide resistant malaria vectors in the field remain susceptible to malathion, despite the presence of Ace1 point mutations. PLoS Genetics, 2022, 18, e1009963.	1.5	12
3	Isolation and Structural Elucidation of Compounds from Pleiocarpa bicarpellata and Their In Vitro Antiprotozoal Activity. Molecules, 2022, 27, 2200.	1.7	4
4	Preparation of new 1,3-dibenzyl tetrahydropyridinylidene ammonium salts and their antimicrobial and anticellular activities. European Journal of Medicinal Chemistry, 2021, 210, 112969.	2.6	6
5	The Alkaloid-Enriched Fraction of Pachysandra terminalis (Buxaceae) Shows Prominent Activity against Trypanosoma brucei rhodesiense. Molecules, 2021, 26, 591.	1.7	4
6	Combination With Tomatidine Improves the Potency of Posaconazole Against Trypanosoma cruzi. Frontiers in Cellular and Infection Microbiology, 2021, 11, 617917.	1.8	6
7	RÃ¼cktitelbild: Antiprotozoische Strukturâ€AktivitÃtsâ€Beziehungen von synthetischen Leucinostatinâ€Derivaten und AufklÃrung ihres Wirkprinzips (Angew. Chem. 28/2021). Angewandte Chemie, 2021, 133, 15792-15792.	1.6	0
8	New Acyl Derivatives of 3-Aminofurazanes and Their Antiplasmodial Activities. Pharmaceuticals, 2021, 14, 412.	1.7	2
9	Antiprotozoal Structureâ€“Activity Relationships of Synthetic Leucinostatin Derivatives and Elucidation of their Mode of Action. Angewandte Chemie - International Edition, 2021, 60, 15613-15621.	7.2	7
10	Salvia officinalis L.: Antitrypanosomal Activity and Active Constituents against Trypanosoma brucei rhodesiense. Molecules, 2021, 26, 3226.	1.7	3
11	Antiprotozoische Strukturâ€AktivitÃtsâ€Beziehungen von synthetischen Leucinostatinâ€Derivaten und AufklÃrung ihres Wirkprinzips. Angewandte Chemie, 2021, 133, 15741-15749.	1.6	0
12	Antiprotozoal Nor-Triterpene Alkaloids from Buxus sempervirens L.. Antibiotics, 2021, 10, 696.	1.5	7
13	Niclosamide Is Active In Vitro against Mycetoma Pathogens. Molecules, 2021, 26, 4005.	1.7	2
14	Boswellic Acids Show In Vitro Activity against Leishmania donovani. Molecules, 2021, 26, 3651.	1.7	6
15	Enantiospecific antitrypanosomal in vitro activity of eflornithine. PLoS Neglected Tropical Diseases, 2021, 15, e0009583.	1.3	3
16	From Magic Bullet to Magic Bomb: Reductive Bioactivation of Antiparasitic Agents. ACS Infectious Diseases, 2021, 7, 2777-2786.	1.8	14
17	8-Amino-6-Methoxyquinolineâ€”Tetrazole Hybrids: Impact of Linkers on Antiplasmodial Activity. Molecules, 2021, 26, 5530.	1.7	3
18	Hygroline derivatives from Schizanthus tricolor and their anti-trypanosomatid and antiplasmodial activities. Phytochemistry, 2021, 192, 112957.	1.4	3

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19	Structure of trypanosome coat protein VSGsur and function in suramin resistance. Nature Microbiology, 2021, 6, 392-400.	5.9	20
20	Unexpected ring-opening of 2,3-dihydropyridines. Monatshefte FÃ¼r Chemie, 2021, 152, 1377-1387.	0.9	0
21	Identification of Antiprotozoal Compounds from BuxusÂsempervirens L. by PLS-Prediction. Molecules, 2021, 26, 6181.	1.7	1
22	Modifications and hybrids of 1,2,3,4-tetrahydropyridinium salts and their antiprotozoal potencies. Monatshefte FÃ¼r Chemie, 2021, 152, 1347-1359.	0.9	1
23	Synthesis and Structure-Activity Relationships of New 2-Phenoxybenzamides with Antiplasmodial Activity. Pharmaceuticals, 2021, 14, 1109.	1.7	1
24	The 3-phosphoinositideâ€“dependent protein kinase 1 is an essential upstream activator of protein kinase A in malaria parasites. PLoS Biology, 2021, 19, e3001483.	2.6	9
25	Structureâ€“Activity Relationship in Pyrazolo[4,3-c]pyridines, First Inhibitors of PEX14â€“PEX5 Proteinâ€“Protein Interaction with Trypanocidal Activity. Journal of Medicinal Chemistry, 2020, 63, 847-879.	2.9	13
26	Use of herbal remedies in the management of sleeping sickness in four northern provinces of Angola. Journal of Ethnopharmacology, 2020, 256, 112382.	2.0	8
27	Pyridine-4(1H)-one Alkaloids from Waltheria indica as Antitrypanosomatid Agents. Journal of Natural Products, 2020, 83, 3363-3371.	1.5	9
28	Non-invasive monitoring of drug action: AÂnew live in vitro assay design for Chagasâ€™ disease drug discovery. PLoS Neglected Tropical Diseases, 2020, 14, e0008487.	1.3	5
29	HPLC-Based Activity Profiling for Antiprotozoal Compounds in Croton gratissimus and Cuscuta hyalina. Frontiers in Pharmacology, 2020, 11, 1246.	1.6	13
30	Mining Sudanese Medicinal Plants for Antiprotozoal Agents. Frontiers in Pharmacology, 2020, 11, 865.	1.6	12
31	Lignans, Amides, and Saponins from Haplophyllum tuberculatum and Their Antiprotozoal Activity. Molecules, 2020, 25, 2825.	1.7	19
32	New Drugs for Human African Trypanosomiasis: A Twenty First Century Success Story. Tropical Medicine and Infectious Disease, 2020, 5, 29.	0.9	83
33	Anti-Trypanosomal Proteasome Inhibitors Cure Hemolymphatic and Meningoencephalic Murine Infection Models of African Trypanosomiasis. Tropical Medicine and Infectious Disease, 2020, 5, 28.	0.9	8
34	100 Years of Suramin. Antimicrobial Agents and Chemotherapy, 2020, 64, .	1.4	121
35	In Vitro Drug Efficacy Testing Against Trypanosoma brucei. Methods in Molecular Biology, 2020, 2116, 781-789.	0.4	0
36	Stochastic Protein Alkylation by Antimalarial Peroxides. ACS Infectious Diseases, 2019, 5, 2067-2075.	1.8	23

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37	New derivatives of 3-azabicyclo[3.2.2]nonanes and their antiprotozoal activities. Monatshefte FÃ¼r Chemie, 2019, 150, 1959-1972.	0.9	1
38	Expression of a specific variant surface glycoprotein has a major impact on suramin sensitivity and endocytosis inTrypanosoma brucei. FASEB BioAdvances, 2019, 1, 595-608.	1.3	12
39	Lysyl-tRNA synthetase as a drug target in malaria and cryptosporidiosis. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 7015-7020.	3.3	94
40	Synthesis of new 1-benzyl tetrahydropyridin-4-ylidene piperidinium salts and their antiplasmodial and antitrypanosomal activities. Medicinal Chemistry Research, 2019, 28, 742-753.	1.1	8
41	Synthesis and structure-activity relationships for new 6-fluoroquinoline derivatives with antiplasmodial activity. Bioorganic and Medicinal Chemistry, 2019, 27, 2052-2065.	1.4	11
42	Anti-malarial ozonides OZ439 and OZ609 tested at clinically relevant compound exposure parameters in a novel ring-stage survival assay. Malaria Journal, 2019, 18, 427.	0.8	13
43	Drug Discovery for Kinetoplastid Diseases: Future Directions. ACS Infectious Diseases, 2019, 5, 152-157.	1.8	78
44	Design, Synthesis, and Biological Evaluation of New 1-(Aryl-1H-pyrrolyl)(phenyl)methyl-1H-imidazole Derivatives as Antiprotozoal Agents. Journal of Medicinal Chemistry, 2019, 62, 1330-1347.	2.9	26
45	Antiprotozoal Activities of Tetrazole-quinolines with Aminopiperidine Linker. Medicinal Chemistry, 2019, 15, 409-416.	0.7	12
46	Isothermal microcalorimetry â€“ A quantitative method to monitor Trypanosoma congolense growth and growth inhibition by trypanocidal drugs in real time. International Journal for Parasitology: Drugs and Drug Resistance, 2018, 8, 159-164.	1.4	13
47	Transporters of Trypanosoma bruceiâ€”phylogeny, physiology, pharmacology. FEBS Journal, 2018, 285, 1012-1023.	2.2	16
48	Beyond immune escape: a variant surface glycoprotein causes suramin resistance in Trypanosoma brucei. Molecular Microbiology, 2018, 107, 57-67.	1.2	26
49	Modifications on tetrahydropyridin-4-ylidene ammonium salts and their antiprotozoal activities. Monatshefte FÃ¼r Chemie, 2018, 149, 801-812.	0.9	3
50	Synthesis of new 1-benzyl tetrahydropyridinylidene ammonium salts and their antimicrobial and anticellular activities. European Journal of Medicinal Chemistry, 2018, 143, 97-106.	2.6	13
51	Come, sweet death: targeting glycosomal protein import for antitrypanosomal drug development. Current Opinion in Microbiology, 2018, 46, 116-122.	2.3	14
52	Host-Microbe Interactions: Parasitology Vol 46. Current Opinion in Microbiology, 2018, 46, vi-viii.	2.3	0
53	Biological evaluation and structure-activity relationships of imidazole-based compounds as antiprotozoal agents. European Journal of Medicinal Chemistry, 2018, 156, 53-60.	2.6	19
54	Cell Penetration, Herbicidal Activity, and inâ€vivoâ€Toxicity of Oligoâ€Arginine Derivatives and of Novel Guanidiniumâ€Rich Compounds Derived from the Biopolymer Cyanophycin. Helvetica Chimica Acta, 2018, 101, e1800112.	1.0	17

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55	Using Yeast Synthetic Lethality To Inform Drug Combination for Malaria. Antimicrobial Agents and Chemotherapy, 2018, 62, .	1.4	2
56	New derivatives of quinoline-4-carboxylic acid with antiplasmodial activity. Bioorganic and Medicinal Chemistry, 2017, 25, 2251-2259.	1.4	6
57	Inhibitors of PEX14 disrupt protein import into glycosomes and kill Trypanosoma parasites. Science, 2017, 355, 1416-1420.	6.0	59
58	New derivatives of 7-chloroquinolin-4-amine with antiprotozoal activity. Bioorganic and Medicinal Chemistry, 2017, 25, 941-948.	1.4	6
59	Comparative sphingolipidomics of disease-causing trypanosomatids reveal unique lifecycle- and taxonomy-specific lipid chemistries. Scientific Reports, 2017, 7, 13617.	1.6	11
60	TbIRK is a signature sequence free potassium channel from Trypanosoma brucei locating to acidocalcisomes. Scientific Reports, 2017, 7, 656.	1.6	13
61	In vitro activity of anti-malarial ozonides against an artemisinin-resistant isolate. Malaria Journal, 2017, 16, 45.	0.8	23
62	Cherchez l'Electron. Molecular Microbiology, 2017, 106, 183-185.	1.2	3
63	Arginine and Lysine Transporters Are Essential for Trypanosoma brucei. PLoS ONE, 2017, 12, e0168775.	1.1	24
64	Aquaglyceroporin-null trypanosomes display glycerol transport defects and respiratory-inhibitor sensitivity. PLoS Pathogens, 2017, 13, e1006307.	2.1	37
65	Drug Resistance in Trypanosoma brucei. , 2017, , 667-676.		0
66	Use of herbal medicine in the management of trypanosomiasis in Angola. Planta Medica International Open, 2017, 4, .	0.3	0
67	Screening of Selected Sudanese Medicinal Plants for In vitro Activity Against Protozoal Neglected Tropical Diseases. , 2017, 4, .		0
68	Identification and characterization of the three members of the CLC family of anion transport proteins in Trypanosoma brucei. PLoS ONE, 2017, 12, e0188219.	1.1	3
69	Assessing anti-T.Âcruzi candidates inÂvitro for sterile cidality. International Journal for Parasitology: Drugs and Drug Resistance, 2016, 6, 165-170.	1.4	38
70	A new approach to chemotherapy: drug-induced differentiation kills African trypanosomes. Scientific Reports, 2016, 6, 22451.	1.6	16
71	Inhibition of Plasmodium falciparum Hsp90 Contributes to the Antimalarial Activities of Aminoalcohol-carbazoles. Journal of Medicinal Chemistry, 2016, 59, 6344-6352.	2.9	34
72	Comparative genomics of drug resistance in Trypanosoma brucei rhodesiense. Cellular and Molecular Life Sciences, 2016, 73, 3387-3400.	2.4	22

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73	Monoclonal Antibodies That Recognize the Alkylation Signature of Antimalarial Ozonides OZ277 (Arterolane) and OZ439 (Artefenomel). ACS Infectious Diseases, 2016, 2, 54-61.	1.8	27
74	An Atypical Mitochondrial Carrier That Mediates Drug Action in Trypanosoma brucei. PLoS Pathogens, 2015, 11, e1004875.	2.1	15
75	A heteromeric potassium channel involved in the modulation of the plasma membrane potential is essential for the survival of African trypanosomes. FASEB Journal, 2015, 29, 3228-3237.	0.2	21
76	Chimerization at the AQP2â€“AQP3 locus is the genetic basis of melarsoprolâ€“pentamidine cross-resistance in clinical Trypanosoma brucei gambiense isolates. International Journal for Parasitology: Drugs and Drug Resistance, 2015, 5, 65-68.	1.4	44
77	Match-making for posaconazole through systems thinking. Trends in Parasitology, 2015, 31, 46-51.	1.5	9
78	TrypanoCyc: a community-led biochemical pathways database for Trypanosoma brucei. Nucleic Acids Research, 2015, 43, D637-D644.	6.5	35
79	Antiprotozoal Activity Profiling of Approved Drugs: A Starting Point toward Drug Repositioning. PLoS ONE, 2015, 10, e0135556.	1.1	81
80	Trypanosoma brucei eflornithine transporter AAT6 is a low-affinity low-selective transporter for neutral amino acids. Biochemical Journal, 2014, 463, 9-18.	1.7	16
81	Comparative Genomics Reveals Multiple Genetic Backgrounds of Human Pathogenicity in the Trypanosoma brucei Complex. Genome Biology and Evolution, 2014, 6, 2811-2819.	1.1	39
82	Trypanosoma brucei adenine-phosphoribosyltransferases mediate adenine salvage and aminopurinol susceptibility but not adenine toxicity. International Journal for Parasitology: Drugs and Drug Resistance, 2014, 4, 55-63.	1.4	26
83	Trypanosoma brucei aquaglyceroporin 2 is a high-affinity transporter for pentamidine and melaminophenyl arsenic drugs and the main genetic determinant of resistance to these drugs. Journal of Antimicrobial Chemotherapy, 2014, 69, 651-663.	1.3	106
84	Differences in Conformational Dynamics between Plasmodium falciparum and Human Hsp90 Orthologues Enable the Structure-Based Discovery of Pathogen-Selective Inhibitors. Journal of Medicinal Chemistry, 2014, 57, 2524-2535.	2.9	38
85	Characterization of choline uptake in Trypanosoma brucei procyclic and bloodstream forms. Molecular and Biochemical Parasitology, 2013, 190, 16-22.	0.5	13
86	Drug resistance in African trypanosomiasis: the melarsoprol and pentamidine story. Trends in Parasitology, 2013, 29, 110-118.	1.5	207
87	Aquaporin 2 Mutations in Trypanosoma brucei gambiense Field Isolates Correlate with Decreased Susceptibility to Pentamidine and Melarsoprol. PLoS Neglected Tropical Diseases, 2013, 7, e2475.	1.3	63
88	In Silico Ionomics Segregates Parasitic from Free-Living Eukaryotes. Genome Biology and Evolution, 2013, 5, 1902-1909.	1.1	4
89	Pyrimidine Salvage in Trypanosoma brucei Bloodstream Forms and the Trypanocidal Action of Halogenated Pyrimidines. Molecular Pharmacology, 2013, 83, 439-453.	1.0	57
90	The genome of the heartworm, Dirofilaria immitis , reveals drug and vaccine targets. FASEB Journal, 2012, 26, 4650-4661.	0.2	124

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91	myo-Inositol Uptake Is Essential for Bulk Inositol Phospholipid but Not Glycosylphosphatidylinositol Synthesis in Trypanosoma brucei. Journal of Biological Chemistry, 2012, 287, 13313-13323.	1.6	34
92	In silico prediction of antimalarial drug target candidates. International Journal for Parasitology: Drugs and Drug Resistance, 2012, 2, 191-199.	1.4	30
93	Antiparasitic agents: new drugs on the horizon. Current Opinion in Pharmacology, 2012, 12, 562-566.	1.7	72
94	The Diamidine Diminazene Aceturate Is a Substrate for the High-Affinity Pentamidine Transporter: Implications for the Development of High Resistance Levels in Trypanosomes. Molecular Pharmacology, 2011, 80, 110-116.	1.0	37
95	Species-specific Typing of DNA Based on Palindrome Frequency Patterns. DNA Research, 2011, 18, 117-124.	1.5	14
96	Genome-Wide Identification of Molecular Mimicry Candidates in Parasites. PLoS ONE, 2011, 6, e17546.	1.1	49
97	A Trk/HKT-Type K ⁺ Transporter from Trypanosoma brucei. Eukaryotic Cell, 2010, 9, 539-546.	3.4	18
98	Adenosine Kinase of T. b. rhodesiense Identified as the Putative Target of 4-[5-(4-phenoxyphenyl)-2H-pyrazol-3-yl]morpholine Using Chemical Proteomics. PLoS Neglected Tropical Diseases, 2009, 3, e506.	1.3	25
99	Genotypic Status of the TbAT1/P2 Adenosine Transporter of Trypanosoma brucei gambiense Isolates from Northwestern Uganda following Melarsoprol Withdrawal. PLoS Neglected Tropical Diseases, 2009, 3, e523.	1.3	16
100	In vitro selection of Haemonchus contortus for benzimidazole resistance reveals a mutation at amino acid 198 of Î²-tubulin. Molecular and Biochemical Parasitology, 2009, 168, 120-122.	0.5	64
101	A new class of anthelmintics effective against drug-resistant nematodes. Nature, 2008, 452, 176-180.	13.7	437
102	Adenosine Kinase Mediates High Affinity Adenosine Salvage in Trypanosoma brucei. Journal of Biological Chemistry, 2008, 283, 5380-5388.	1.6	40
103	Loss of the High-Affinity Pentamidine Transporter Is Responsible for High Levels of Cross-Resistance between Arsenical and Diamidine Drugs in African Trypanosomes. Molecular Pharmacology, 2007, 71, 1098-1108.	1.0	113
104	Chemotherapeutic Strategies Against Trypanosoma brucei: Drug Targets vs. Drug Targeting. Current Pharmaceutical Design, 2007, 13, 555-567.	0.9	72
105	Adenosine Kinase ofTrypanosoma bruceiand Its Role in Susceptibility to Adenosine Antimetabolites. Antimicrobial Agents and Chemotherapy, 2007, 51, 3895-3901.	1.4	37
106	Genotypic and phenotypic characterization of Trypanosoma brucei gambiense isolates from Ibba, South Sudan, an area of high melarsoprol treatment failure rate. Acta Tropica, 2007, 104, 84-90.	0.9	39
107	Phenotyping and genotyping of Haemonchus contortus isolates reveals a new putative candidate mutation for benzimidazole resistance in nematodes. Veterinary Parasitology, 2007, 144, 313-320.	0.7	223
108	Melarsoprol- and pentamidine-resistant Trypanosoma brucei rhodesiense populations and their cross-resistance. International Journal for Parasitology, 2007, 37, 1443-1448.	1.3	30

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109	Combined contribution of TbAT1 and TbMRPA to drug resistance in Trypanosoma brucei. Molecular and Biochemical Parasitology, 2006, 150, 364-366.	0.5	21
110	Identification of GPI anchor attachment signals by a Kohonen self-organizing map. Bioinformatics, 2005, 21, 1846-1852.	1.8	275
111	Molecular Pharmacology of Adenosine Transport in Trypanosoma brucei: P1/P2 Revisited. Molecular Pharmacology, 2005, 68, 589-595.	1.0	49
112	Drug transport and drug resistance in African trypanosomes. Drug Resistance Updates, 2003, 6, 281-290.	6.5	60
113	Mechanisms of Arsenical and Diamidine Uptake and Resistance in Trypanosoma brucei. Eukaryotic Cell, 2003, 2, 1003-1008.	3.4	186
114	An anti-contamination cocktail for the in vitro isolation and cultivation of parasitic protozoa. Parasitology Research, 2002, 88, 172-174.	0.6	38
115	Identification and characterization of trypanocides by functional expression of an adenosine transporter from Trypanosoma brucei in yeast. Journal of Molecular Medicine, 2001, 79, 121-127.	1.7	27
116	Genetic variants of the TbAT1 adenosine transporter from African trypanosomes in relapse infections following melarsoprol therapy. Molecular and Biochemical Parasitology, 2001, 117, 73-81.	0.5	81
117	A Nucleoside Transporter from Trypanosoma brucei Involved in Drug Resistance. Science, 1999, 285, 242-244.	6.0	245
118	Pharmacokinetic profiles reconcile in vitro and in vivo activities of novel trypanocidal compounds. Matters, 0, , .	1.0	0