

Michael Delves

List of Publications by Year in descending order

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Version: 2024-02-01

52
papers

3,892
citations

136950

32
h-index

175258

52
g-index

55
all docs

55
docs citations

55
times ranked

4384
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification and Profiling of a Novel Diazaspiro[3.4]octane Chemical Series Active against Multiple Stages of the Human Malaria Parasite <i>Plasmodium falciparum</i> and Optimization Efforts. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 2291-2309.	6.4	11
2	Inhibition of Resistance-Refractory <i>P. falciparum</i> Kinase PKG Delivers Prophylactic, Blood Stage, and Transmission-Blocking Antiplasmodial Activity. <i>Cell Chemical Biology</i> , 2020, 27, 806-816.e8.	5.2	56
3	Fueling Open Innovation for Malaria Transmission-Blocking Drugs: Hundreds of Molecules Targeting Early Parasite Mosquito Stages. <i>Frontiers in Microbiology</i> , 2019, 10, 2134.	3.5	31
4	Substituted Aminoacetamides as Novel Leads for Malaria Treatment. <i>ChemMedChem</i> , 2019, 14, 1329-1335.	3.2	5
5	Lysyl-tRNA synthetase as a drug target in malaria and cryptosporidiosis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 7015-7020.	7.1	94
6	The antimalarial screening landscape—looking beyond the asexual blood stage. <i>Current Opinion in Chemical Biology</i> , 2019, 50, 1-9.	6.1	27
7	Transmission-blocking Effects of Primaquine and Methylene Blue Suggest <i>Plasmodium falciparum</i> Gametocyte Sterilization Rather Than Effects on Sex Ratio. <i>Clinical Infectious Diseases</i> , 2019, 69, 1436-1439.	5.8	21
8	8-Aminoquinolines with an Aminoxyalkyl Side Chain Exert in vitro Dual-Stage Antiplasmodial Activity. <i>ChemMedChem</i> , 2019, 14, 501-511.	3.2	6
9	Kinesin-8B controls basal body function and flagellum formation and is key to malaria transmission. <i>Life Science Alliance</i> , 2019, 2, e201900488.	2.8	33
10	Polyamidoamine Nanoparticles for the Oral Administration of Antimalarial Drugs. <i>Pharmaceutics</i> , 2018, 10, 225.	4.5	17
11	Allele-Specific Isothermal Amplification Method Using Unmodified Self-Stabilizing Competitive Primers. <i>Analytical Chemistry</i> , 2018, 90, 11972-11980.	6.5	22
12	A high throughput screen for next-generation leads targeting malaria parasite transmission. <i>Nature Communications</i> , 2018, 9, 3805.	12.8	92
13	An inexpensive open source 3D-printed membrane feeder for human malaria transmission studies. <i>Malaria Journal</i> , 2018, 17, 282.	2.3	13
14	Antimalarial Transmission-Blocking Interventions: Past, Present, and Future. <i>Trends in Parasitology</i> , 2018, 34, 735-746.	3.3	73
15	Antimalarial efficacy of MMV390048, an inhibitor of <i>Plasmodium</i> phosphatidylinositol 4-kinase. <i>Science Translational Medicine</i> , 2017, 9, .	12.4	204
16	Hundreds of dual-stage antimalarial molecules discovered by a functional gametocyte screen. <i>Nature Communications</i> , 2017, 8, 15160.	12.8	42
17	A tetraoxane-based antimalarial drug candidate that overcomes PfK13-C580Y dependent artemisinin resistance. <i>Nature Communications</i> , 2017, 8, 15159.	12.8	51
18	3-Hydroxy-N ² -arylidenepropanehydrazonamides with Halo-Substituted Phenanthrene Scaffolds Cure <i>P. berghei</i> Infected Mice When Administered Perorally. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 6036-6044.	6.4	4

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19	Adaptation of targeted nanocarriers to changing requirements in antimalarial drug delivery. <i>Nanomedicine: Nanotechnology, Biology, and Medicine</i> , 2017, 13, 515-525.	3.3	49
20	Hexahydroquinolines are antimalarial candidates with potent blood-stage and transmission-blocking activity. <i>Nature Microbiology</i> , 2017, 2, 1403-1414.	13.3	47
21	Open Source Drug Discovery with the Malaria Box Compound Collection for Neglected Diseases and Beyond. <i>PLoS Pathogens</i> , 2016, 12, e1005763.	4.7	244
22	Characterization of Novel Antimalarial Compound ACT-451840: Preclinical Assessment of Activity and Dose-Response Efficacy Modeling. <i>PLoS Medicine</i> , 2016, 13, e1002138.	8.4	35
23	Discovery of a Quinoline-4-carboxamide Derivative with a Novel Mechanism of Action, Multistage Antimalarial Activity, and Potent in Vivo Efficacy. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 9672-9685.	6.4	66
24	Routine in vitro culture of <i>P. falciparum</i> gametocytes to evaluate novel transmission-blocking interventions. <i>Nature Protocols</i> , 2016, 11, 1668-1680.	12.0	115
25	Open Source Drug Discovery: Highly Potent Antimalarial Compounds Derived from the Tres Cantos Arylpyrroles. <i>ACS Central Science</i> , 2016, 2, 687-701.	11.3	68
26	A Malaria Transmission-Blocking (+)-Usnic Acid Derivative Prevents <i>Plasmodium</i> Zygote-to-Ookinete Maturation in the Mosquito Midgut. <i>ACS Chemical Biology</i> , 2016, 11, 3461-3472.	3.4	13
27	Use of <i>Plasmodium falciparum</i> culture-adapted field isolates for in vitro exflagellation-blocking assay. <i>Malaria Journal</i> , 2015, 14, 234.	2.3	10
28	Lead Clinical and Preclinical Antimalarial Drugs Can Significantly Reduce Sporozoite Transmission to Vertebrate Populations. <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 490-497.	3.2	23
29	Histone Methyltransferase Inhibitors Are Orally Bioavailable, Fast-Acting Molecules with Activity against Different Species Causing Malaria in Humans. <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 950-959.	3.2	43
30	A novel multiple-stage antimalarial agent that inhibits protein synthesis. <i>Nature</i> , 2015, 522, 315-320.	27.8	353
31	The <i>Plasmodium berghei</i> sexual stage antigen PSOP12 induces anti-malarial transmission blocking immunity both in vivo and in vitro. <i>Vaccine</i> , 2015, 33, 437-445.	3.8	30
32	A long-duration dihydroorotate dehydrogenase inhibitor (DSM265) for prevention and treatment of malaria. <i>Science Translational Medicine</i> , 2015, 7, 296ra111.	12.4	254
33	Imaging-Based High-Throughput Screening Assay To Identify New Molecules with Transmission-Blocking Potential against <i>Plasmodium falciparum</i> Female Gamete Formation. <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 3298-3305.	3.2	45
34	An essential role of the basal body protein SAS in <i>Plasmodium</i> male gamete development and malaria transmission. <i>Cellular Microbiology</i> , 2015, 17, 191-206.	2.1	37
35	Changes in metabolic phenotypes of <i>Plasmodium falciparum</i> in vitro cultures during gametocyte development. <i>Malaria Journal</i> , 2014, 13, 468.	2.3	36
36	Assessment of therapeutic responses to gametocytocidal drugs in <i>Plasmodium falciparum</i> malaria. <i>Malaria Journal</i> , 2014, 13, 483.	2.3	61

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37	A Male and Female Gametocyte Functional Viability Assay To Identify Biologically Relevant Malaria Transmission-Blocking Drugs. <i>Antimicrobial Agents and Chemotherapy</i> , 2014, 58, 7292-7302.	3.2	112
38	Pyrazoleamide compounds are potent antimalarials that target Na ⁺ homeostasis in intraerythrocytic <i>Plasmodium falciparum</i> . <i>Nature Communications</i> , 2014, 5, 5521.	12.8	108
39	Quantitative analysis of <i>Plasmodium</i> ookinete motion in three dimensions suggests a critical role for cell shape in the biomechanics of malaria parasite gliding motility. <i>Cellular Microbiology</i> , 2014, 16, 734-750.	2.1	45
40	Quinolone-3-Diarylethers: A New Class of Antimalarial Drug. <i>Science Translational Medicine</i> , 2013, 5, 177ra37.	12.4	187
41	Male and Female <i>Plasmodium falciparum</i> Mature Gametocytes Show Different Responses to Antimalarial Drugs. <i>Antimicrobial Agents and Chemotherapy</i> , 2013, 57, 3268-3274.	3.2	158
42	<i>Plasmodium</i> cell biology should inform strategies used in the development of antimalarial transmission-blocking drugs. <i>Future Medicinal Chemistry</i> , 2012, 4, 2251-2263.	2.3	27
43	The design and interpretation of laboratory assays measuring mosquito transmission of <i>Plasmodium</i> . <i>Trends in Parasitology</i> , 2012, 28, 457-465.	3.3	32
44	A high-throughput assay for the identification of malarial transmission-blocking drugs and vaccines. <i>International Journal for Parasitology</i> , 2012, 42, 999-1006.	3.1	59
45	Measuring the blockade of malaria transmission – An analysis of the Standard Membrane Feeding Assay. <i>International Journal for Parasitology</i> , 2012, 42, 1037-1044.	3.1	162
46	Spatial Localisation of Actin Filaments across Developmental Stages of the Malaria Parasite. <i>PLoS ONE</i> , 2012, 7, e32188.	2.5	69
47	Generation of quinolone antimalarials targeting the <i>Plasmodium falciparum</i> mitochondrial respiratory chain for the treatment and prophylaxis of malaria. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 8298-8303.	7.1	143
48	A GFP-Actin reporter line to explore microfilament dynamics across the malaria parasite lifecycle. <i>Molecular and Biochemical Parasitology</i> , 2012, 182, 93-96.	1.1	15
49	The Activities of Current Antimalarial Drugs on the Life Cycle Stages of <i>Plasmodium</i> : A Comparative Study with Human and Rodent Parasites. <i>PLoS Medicine</i> , 2012, 9, e1001169.	8.4	301
50	Use of a Selective Inhibitor To Define the Chemotherapeutic Potential of the Plasmodial Hexose Transporter in Different Stages of the Parasite's Life Cycle. <i>Antimicrobial Agents and Chemotherapy</i> , 2011, 55, 2824-2830.	3.2	39
51	A semi-automated method for counting fluorescent malaria oocysts increases the throughput of transmission blocking studies. <i>Malaria Journal</i> , 2010, 9, 35.	2.3	45
52	<i>Plasmodium</i> male development gene-1 (mdv-1) is important for female sexual development and identifies a polarised plasma membrane during zygote development. <i>International Journal for Parasitology</i> , 2009, 39, 755-761.	3.1	36