

Vicky M. AVERY

List of Publications by Year in descending order

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Version: 2024-02-01

219
papers

8,970
citations

44069

48
h-index

60623

81
g-index

239
all docs

239
docs citations

239
times ranked

10412
citing authors

#	ARTICLE	IF	CITATIONS
1	Assay development in leishmaniasis drug discovery: a comprehensive review. <i>Expert Opinion on Drug Discovery</i> , 2022, 17, 151-166.	5.0	7
2	Isolation of Antimalarial Agents From Indonesian Medicinal Plants: <i>Swietenia mahagoni</i> and <i>Pluchea indica</i> . <i>Natural Product Communications</i> , 2022, 17, 1934578X2110689.	0.5	1
3	Î±-Synuclein Aggregation Inhibitory Prunolides and a Dibrominated Î²-Carboline Sulfamate from the Ascidian <i>Sycoicum prunum</i> . <i>Journal of Natural Products</i> , 2022, 85, 441-452.	3.0	8
4	Investigating the antiplasmodial activity of substituted cyclopentadienyl rhodium and iridium complexes of 2-(2-pyridyl)benzimidazole. <i>Journal of Organometallic Chemistry</i> , 2022, 962, 122273.	1.8	10
5	Temporal and Wash-Out Studies Identify Medicines for Malaria Venture Pathogen Box Compounds with Fast-Acting Activity against Both <i>Trypanosoma cruzi</i> and <i>Trypanosoma brucei</i> . <i>Microorganisms</i> , 2022, 10, 1287.	3.6	2
6	Addressing the tumour microenvironment in early drug discovery: a strategy to overcome drug resistance and identify novel targets for cancer therapy. <i>Drug Discovery Today</i> , 2021, 26, 663-676.	6.4	22
7	Tedaniophorbins A and B: Novel Fluorescent Pteridine Alkaloids Incorporating a Thiomorpholine from the Sponge <i>Tedaniophora ceratosis</i> . <i>Marine Drugs</i> , 2021, 19, 95.	4.6	8
8	Synthesis and antimicrobial study of organoiridium amido-sulfadoxine complexes. <i>Inorganica Chimica Acta</i> , 2021, 517, 120175.	2.4	8
9	Structure activity refinement of phenylsulfonyl piperazines as antimalarials that block erythrocytic invasion. <i>European Journal of Medicinal Chemistry</i> , 2021, 214, 113253.	5.5	11
10	Discovery of Potent and Fast-Acting Antimalarial Bis-1,2,4-triazines. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 4150-4162.	6.4	14
11	Repositioning and Characterization of 1-(Pyridin-4-yl)pyrrolidin-2-one Derivatives as <i>Plasmodium</i> Cytoplasmic Prolyl-tRNA Synthetase Inhibitors. <i>ACS Infectious Diseases</i> , 2021, 7, 1680-1689.	3.8	14
12	Synthesis of New Triazolopyrazine Antimalarial Compounds. <i>Molecules</i> , 2021, 26, 2421.	3.8	3
13	Synthesis and Evaluation of the Tetracyclic Ring-System of Isocryptolepine and Regioisomers for Antimalarial, Antiproliferative and Antimicrobial Activities. <i>Molecules</i> , 2021, 26, 3268.	3.8	7
14	Bioactive half-sandwich Rh and Ir bipyridyl complexes containing artemisinin. <i>Journal of Inorganic Biochemistry</i> , 2021, 219, 111408.	3.5	7
15	Abstract 1130: First-in-class KAT6A/KAT6B inhibitor CTx-648 (PF-9363) demonstrates potent anti-tumor activity in ER+ breast cancer with KAT6A dysregulation. <i>Cancer Research</i> , 2021, 81, 1130-1130.	0.9	8
16	The Novel bis-1,2,4-Triazine MIPS-0004373 Demonstrates Rapid and Potent Activity against All Blood Stages of the Malaria Parasite. <i>Antimicrobial Agents and Chemotherapy</i> , 2021, 65, e0031121.	3.2	4
17	Discovery and Structure-Activity Relationships of Quinazolinone-2-carboxamide Derivatives as Novel Orally Efficacious Antimalarials. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 12582-12602.	6.4	11
18	Property activity refinement of 2-anilino 4-amino substituted quinazolines as antimalarials with fast acting asexual parasite activity. <i>Bioorganic Chemistry</i> , 2021, 117, 105359.	4.1	8

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19	Discovery and development of 2-aminobenzimidazoles as potent antimalarials. <i>European Journal of Medicinal Chemistry</i> , 2021, 221, 113518.	5.5	11
20	Discovery of Potent <i>N</i> -Ethylurea Pyrazole Derivatives as Dual Inhibitors of <i>Trypanosoma brucei</i> and <i>Trypanosoma cruzi</i> . <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 278-285.	2.8	15
21	Antiplasmodial Bis-Indole Alkaloids from the Bark of <i>Flindersia pimenteliana</i> . <i>Planta Medica</i> , 2020, 86, 19-25.	1.3	6
22	HBO1 is required for the maintenance of leukaemia stem cells. <i>Nature</i> , 2020, 577, 266-270.	27.8	105
23	Re-evaluating pretomanid analogues for Chagas disease: Hit-to-lead studies reveal both <i>in vitro</i> and <i>in vivo</i> trypanocidal efficacy. <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112849.	5.5	13
24	Citronamine A, an Antiplasmodial Isoquinoline Alkaloid from the Australian Marine Sponge <i>Citronia astra</i> . <i>Organic Letters</i> , 2020, 22, 9574-9578.	4.6	8
25	Hemin Prevents Increased Glycolysis in Macrophages upon Activation: Protection by Microbiota-Derived Metabolites of Polyphenols. <i>Antioxidants</i> , 2020, 9, 1109.	5.1	8
26	Prenylated Flavonoids from the Roots of <i>Tephrosia rhodesica</i> . <i>Journal of Natural Products</i> , 2020, 83, 2390-2398.	3.0	6
27	Investigation of thiazolylbenzothiophenamides as potential agents for African sleeping sickness. <i>RSC Medicinal Chemistry</i> , 2020, 11, 1413-1422.	3.9	2
28	Antiplasmodial Alkaloids from the Australian Bryozoan <i>Amathia lamourouxi</i> . <i>Journal of Natural Products</i> , 2020, 83, 3435-3444.	3.0	12
29	Secoiridoids and Iridoids from <i>Morinda asteroscepa</i> . <i>Journal of Natural Products</i> , 2020, 83, 2641-2646.	3.0	7
30	Metabolic Roles of Androgen Receptor and Tip60 in Androgen-Dependent Prostate Cancer. <i>International Journal of Molecular Sciences</i> , 2020, 21, 6622.	4.1	9
31	Antitubercular and Antiparasitic 2-Nitroimidazopyrazinones with Improved Potency and Solubility. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 15726-15751.	6.4	17
32	Hit-to-lead optimization of novel benzimidazole phenylacetamides as broad spectrum trypanosomacides. <i>RSC Medicinal Chemistry</i> , 2020, 11, 685-695.	3.9	5
33	Investigation of pyrimidine nucleoside analogues as chemical probes to assess compound effects on the proliferation of <i>Trypanosoma cruzi</i> intracellular parasites. <i>PLoS Neglected Tropical Diseases</i> , 2020, 14, e0008068.	3.0	10
34	A Meroisoprenoid, Heptenolides, and <i>C</i> -Benzylated Flavonoids from <i>Sphaerocoryne gracilis</i> ssp. <i>gracilis</i> . <i>Journal of Natural Products</i> , 2020, 83, 316-322.	3.0	12
35	Orthoscuticellines \hat{E} , \hat{I}^2 -Carboline Alkaloids from the Bryozoan <i>Orthoscuticella ventricosa</i> Collected in Australia. <i>Journal of Natural Products</i> , 2020, 83, 422-428.	3.0	27
36	A <i>Plasmodium vivax</i> experimental human infection model for evaluating efficacy of interventions. <i>Journal of Clinical Investigation</i> , 2020, 130, 2920-2927.	8.2	25

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37	Antiplasmodial, Antimicrobial and Cytotoxic Activities of Extracts from Selected Medicinal Plants Growing in Tanzania. <i>Journal of Biologically Active Products From Nature</i> , 2020, 10, 165-176.	0.3	1
38	Desymmetrization Reactions of Indigo with Grignard Reagents for the Synthesis of Selective Antiplasmodial [1 <i>H</i>]-3- ² -Aryl-2,2-diindol-3-ones. <i>Journal of Organic Chemistry</i> , 2019, 84, 3.2 11228-11239.		6
39	Total Synthesis of the Antimalarial Ascidian Natural Product Albopunctatone. <i>Organic Letters</i> , 2019, 21, 5519-5523.	4.6	7
40	A New Benzopyranyl Cadenane Sesquiterpene and Other Antiplasmodial and Cytotoxic Metabolites from <i>Cleistochlamys kirkii</i> . <i>Molecules</i> , 2019, 24, 2746.	3.8	14
41	Sulfide, sulfoxide and sulfone bridged acyclic nucleoside phosphonates as inhibitors of the <i>Plasmodium falciparum</i> and human 6-oxopurine phosphoribosyltransferases: Synthesis and evaluation. <i>European Journal of Medicinal Chemistry</i> , 2019, 183, 111667.	5.5	12
42	The Molecular Effects of Sulforaphane and Capsaicin on Metabolism upon Androgen and Tip60 Activation of Androgen Receptor. <i>International Journal of Molecular Sciences</i> , 2019, 20, 5384.	4.1	15
43	The cubane paradigm in bioactive molecule discovery: further scope, limitations and the cyclooctatetraene complement. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 6790-6798.	2.8	49
44	Substituted Aminoacetamides as Novel Leads for Malaria Treatment. <i>ChemMedChem</i> , 2019, 14, 1329-1335.	3.2	5
45	Acrotrione: An Oxidized Xanthene from the Roots of <i>Acronychia pubescens</i> . <i>Journal of Natural Products</i> , 2019, 82, 1019-1023.	3.0	13
46	3,3-Disubstituted 5-Bi(1,2,4-triazine) Derivatives with Potent in Vitro and in Vivo Antimalarial Activity. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 2485-2498.	6.4	16
47	Inhibition of Plasmeprin V Activity Blocks <i>Plasmodium falciparum</i> Gametocytogenesis and Transmission to Mosquitoes. <i>Cell Reports</i> , 2019, 29, 3796-3806.e4.	6.4	25
48	8-Aminoquinolines with an Aminoxyalkyl Side Chain Exert in vitro Dual-Stage Antiplasmodial Activity. <i>ChemMedChem</i> , 2019, 14, 501-511.	3.2	6
49	Hydroxamic Acid Inhibitors Provide Cross-Species Inhibition of <i>Plasmodium</i> M1 and M17 Aminopeptidases. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 622-640.	6.4	30
50	HSQC-TOCSY Fingerprinting for Prioritization of Polyketide- and Peptide-Producing Microbial Isolates. <i>Journal of Natural Products</i> , 2018, 81, 957-965.	3.0	23
51	Organometallic Conjugates of the Drug Sulfadoxine for Combatting Antimicrobial Resistance. <i>Chemistry - A European Journal</i> , 2018, 24, 10078-10090.	3.3	28
52	3-pyridyl inhibitors with novel activity against <i>Trypanosoma cruzi</i> reveal in vitro profiles can aid prediction of putative cytochrome P450 inhibition. <i>Scientific Reports</i> , 2018, 8, 4901.	3.3	19
53	Microthecaline A, a Quinoline Serrulatane Alkaloid from the Roots of the Australian Desert Plant <i>Eremophila microtheca</i> . <i>Journal of Natural Products</i> , 2018, 81, 1079-1083.	3.0	33
54	Design, Synthesis, and Biological Evaluation of 2-Nitroimidazopyrazin-one/-es with Antitubercular and Antiparasitic Activity. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 11349-11371.	6.4	22

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55	Î²-Triketoneâ€“Monoterpene Hybrids from the Flowers of the Australian Tree <i>Corymbia intermedia</i> . Journal of Natural Products, 2018, 81, 2455-2461.	3.0	8
56	Target Validation and Identification of Novel Boronate Inhibitors of the <i>Plasmodium falciparum</i> Proteasome. Journal of Medicinal Chemistry, 2018, 61, 10053-10066.	6.4	54
57	One-pot, multi-component synthesis and structure-activity relationships of peptoid-based histone deacetylase (HDAC) inhibitors targeting malaria parasites. European Journal of Medicinal Chemistry, 2018, 158, 801-813.	5.5	29
58	Routine In Vitro Culture of <i>Plasmodium falciparum</i> : Experimental Consequences?. Trends in Parasitology, 2018, 34, 564-575.	3.3	17
59	Antiplasmodial Î²-Triketoneâ€“Flavanone Hybrids from the Flowers of the Australian Tree <i>Corymbia torelliana</i> . Journal of Natural Products, 2018, 81, 1588-1597.	3.0	16
60	Cascade reactions of indigo with oxiranes and aziridines: efficient access to dihydropyrazinodiindoles and spiro-oxazocinodiindoles. Organic and Biomolecular Chemistry, 2018, 16, 6006-6016.	2.8	12
61	HSQC-TOCSY Fingerprinting-Directed Discovery of Antiplasmodial Polyketides from the Marine Ascidian-Derived <i>Streptomyces</i> sp. (USC-16018). Marine Drugs, 2018, 16, 189.	4.6	17
62	Frontispiece: Organometallic Conjugates of the Drug Sulfadoxine for Combatting Antimicrobial Resistance. Chemistry - A European Journal, 2018, 24, .	3.3	0
63	Doxorubicin resistance in breast cancer cells is mediated by extracellular matrix proteins. BMC Cancer, 2018, 18, 41.	2.6	234
64	<i>Plasmodium falciparum</i> In Vitro Culture â€“ The Highs and Lows. Trends in Parasitology, 2018, 34, 812-813.	3.3	2
65	SC83288 is a clinical development candidate for the treatment of severe malaria. Nature Communications, 2017, 8, 14193.	12.8	19
66	Optimization of 2-Anilino 4-Amino Substituted Quinazolines into Potent Antimalarial Agents with Oral in Vivo Activity. Journal of Medicinal Chemistry, 2017, 60, 1171-1188.	6.4	43
67	The need to compare: assessing the level of agreement of three high-throughput assays against <i>Plasmodium falciparum</i> mature gametocytes. Scientific Reports, 2017, 7, 45992.	3.3	15
68	Leishmaniasis drug discovery: recent progress and challenges in assay development. Drug Discovery Today, 2017, 22, 1516-1531.	6.4	145
69	Antiplasmodial Î²-triketones from the flowers of the Australian tree <i>Angophora woodsiana</i> . Bioorganic and Medicinal Chemistry Letters, 2017, 27, 2602-2607.	2.2	13
70	Polyoxygenated Cyclohexenes and Other Constituents of <i>Cleistochlamys kirkii</i> Leaves. Journal of Natural Products, 2017, 80, 114-125.	3.0	27
71	Pterocarpan and isoflavones from the root bark of <i>Millettia micans</i> and of <i>Millettia dura</i> . Phytochemistry Letters, 2017, 21, 216-220.	1.2	12
72	<i>Plasmodium falciparum</i> in vitro continuous culture conditions: A comparison of parasite susceptibility and tolerance to anti-malarial drugs throughout the asexual intra-erythrocytic life cycle. International Journal for Parasitology: Drugs and Drug Resistance, 2017, 7, 295-302.	3.4	24

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73	Synthesis of antimalarial amide analogues based on the plant serrulatane diterpenoid 3,7,8-trihydroxyserrulat-14-en-19-oic acid. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 4091-4095.	2.2	14
74	Design and Synthesis of Terephthalic Acid-Based Histone Deacetylase Inhibitors with Dual-Stage Anti-Plasmodium Activity. <i>ChemMedChem</i> , 2017, 12, 1627-1636.	3.2	14
75	Pimentelamines A-C, Indole Alkaloids Isolated from the Leaves of the Australian Tree <i>Flindersia pimenteliana</i> . <i>Journal of Natural Products</i> , 2017, 80, 3211-3217.	3.0	27
76	Screening the Medicines for Malaria Venture Pathogen Box across Multiple Pathogens Reclassifies Starting Points for Open-Source Drug Discovery. <i>Antimicrobial Agents and Chemotherapy</i> , 2017, 61, .	3.2	106
77	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
78	Antiplasmodial dihetarylthioethers target the coenzyme A synthesis pathway in <i>Plasmodium falciparum</i> erythrocytic stages. <i>Malaria Journal</i> , 2017, 16, 192.	2.3	13
79	Three Chalconoids and a Pterocarpene from the Roots of <i>Tephrosia aequilata</i> . <i>Molecules</i> , 2017, 22, 318.	3.8	11
80	Screening a Natural Product-Based Library against Kinetoplastid Parasites. <i>Molecules</i> , 2017, 22, 1715.	3.8	53
81	Hexahydroquinolines are antimalarial candidates with potent blood-stage and transmission-blocking activity. <i>Nature Microbiology</i> , 2017, 2, 1403-1414.	13.3	47
82	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
83	Metabolomics and lipidomics reveal perturbation of sphingolipid metabolism by a novel anti-trypanosomal 3-(oxazolo[4,5-b]pyridine-2-yl)anilide. <i>Metabolomics</i> , 2016, 12, 1.	3.0	28
84	Innovative in vitro models for breast cancer drug discovery. <i>Drug Discovery Today: Disease Models</i> , 2016, 21, 11-16.	1.2	3
85	Naseseazine C, a new anti-plasmodial dimeric diketopiperazine from a marine sediment derived <i>Streptomyces</i> sp.. <i>Tetrahedron Letters</i> , 2016, 57, 5893-5895.	1.4	32
86	An evaluation of Minor Groove Binders as anti- <i>Trypanosoma brucei brucei</i> therapeutics. <i>European Journal of Medicinal Chemistry</i> , 2016, 116, 116-125.	5.5	24
87	Large-scale production of <i>Plasmodium falciparum</i> gametocytes for malaria drug discovery. <i>Nature Protocols</i> , 2016, 11, 976-992.	12.0	49
88	Selective anti-malarial minor groove binders. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 3326-3329.	2.2	13
89	Diversity-oriented synthesis yields novel multistage antimalarial inhibitors. <i>Nature</i> , 2016, 538, 344-349.	27.8	214
90	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

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91	Expression of the thioredoxin system in an in vivo-like cancer cell environment upon auranofin treatment. <i>European Journal of Cell Biology</i> , 2016, 95, 378-388.	3.6	4
92	Cancer drug discovery: recent innovative approaches to tumor modeling. <i>Expert Opinion on Drug Discovery</i> , 2016, 11, 885-894.	5.0	23
93	Hit-to-Lead Optimization of a Novel Class of Potent, Broad-Spectrum Trypanosomacides. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 9686-9720.	6.4	30
94	Assessing Drug Efficacy in a Miniaturized Pancreatic Cancer <i>In Vitro</i> 3D Cell Culture Model. <i>Assay and Drug Development Technologies</i> , 2016, 14, 367-380.	1.2	14
95	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
96	Development of ethynyl-2â€²-deoxyuridine chemical probes for cell proliferation. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 4272-4280.	3.0	6
97	Biological characterization of chemically diverse compounds targeting the Plasmodium falciparum coenzyme A synthesis pathway. <i>Parasites and Vectors</i> , 2016, 9, 589.	2.5	16
98	Trisubstituted Pyrimidines as Efficacious and Fast-Acting Antimalarials. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 6101-6120.	6.4	13
99	Potent dual inhibitors of Plasmodium falciparum M1 and M17 aminopeptidases through optimization of S1 pocket interactions. <i>European Journal of Medicinal Chemistry</i> , 2016, 110, 43-64.	5.5	46
100	Luciferase-Based, High-Throughput Assay for Screening and Profiling Transmission-Blocking Compounds against Plasmodium falciparum Gametocytes. <i>Antimicrobial Agents and Chemotherapy</i> , 2016, 60, 2097-2107.	3.2	62
101	Identification and Characterization of FTY720 for the Treatment of Human African Trypanosomiasis. <i>Antimicrobial Agents and Chemotherapy</i> , 2016, 60, 1859-1861.	3.2	5
102	A novel class of indole alkaloids isolated from Flindersia pimenteliana (Rutaceae). <i>Planta Medica</i> , 2016, 81, S1-S381.	1.3	0
103	Future treatment options for human African trypanosomiasis. <i>Expert Review of Anti-Infective Therapy</i> , 2015, 13, 1429-1432.	4.4	10
104	Development and application of a sensitive, phenotypic, high-throughput image-based assay to identify compound activity against Trypanosoma cruzi amastigotes. <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2015, 5, 215-228.	3.4	39
105	A simple and predictive phenotypic High Content Imaging assay for Plasmodium falciparum mature gametocytes to identify malaria transmission blocking compounds. <i>Scientific Reports</i> , 2015, 5, 16414.	3.3	46
106	Splenic Retention of Plasmodium falciparum Gametocytes To Block the Transmission of Malaria. <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 4206-4214.	3.2	24
107	Rotenoids, Flavonoids, and Chalcones from the Root Bark of <i>Millettia usaramensis</i> . <i>Journal of Natural Products</i> , 2015, 78, 2932-2939.	3.0	33
108	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

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109	A novel multiple-stage antimalarial agent that inhibits protein synthesis. <i>Nature</i> , 2015, 522, 315-320.	27.8	353
110	6-Arylpyrazine-2-carboxamides: A New Core for <i>Trypanosoma brucei</i> Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 6753-6765.	6.4	18
111	Mechanical clearance of red blood cells by the human spleen: Potential therapeutic applications of a biomimetic RBC filtration method. <i>Transfusion Clinique Et Biologique</i> , 2015, 22, 151-157.	0.4	33
112	Profiling the anti-protozoal activity of anti-cancer HDAC inhibitors against <i>Plasmodium</i> and <i>Trypanosoma</i> parasites. <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2015, 5, 117-126.	3.4	77
113	A long-duration dihydroorotate dehydrogenase inhibitor (DSM265) for prevention and treatment of malaria. <i>Science Translational Medicine</i> , 2015, 7, 296ra111.	12.4	254
114	Design and Synthesis of a Screening Library Using the Natural Product Scaffold 3-Chloro-4-hydroxyphenylacetic Acid. <i>Journal of Natural Products</i> , 2015, 78, 914-918.	3.0	10
115	Evaluation of chemotherapeutics in a three-dimensional breast cancer model. <i>Journal of Cancer Research and Clinical Oncology</i> , 2015, 141, 951-959.	2.5	67
116	The synthesis, antimalarial activity and CoMFA analysis of novel aminoalkylated quercetin analogs. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 327-332.	2.2	36
117	Synthesis and evaluation of phenoxymethylbenzamide analogues as anti-trypanosomal agents. <i>MedChemComm</i> , 2015, 6, 403-406.	3.4	6
118	Advanced Cell Culture Techniques for Cancer Drug Discovery. <i>Biology</i> , 2014, 3, 345-367.	2.8	210
119	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
120	(+)-SJ733, a clinical candidate for malaria that acts through ATP4 to induce rapid host-mediated clearance of <i>Plasmodium</i> . <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, E5455-62.	7.1	199
121	Synthesis, Antimalarial Properties, and SAR Studies of Alkoxyurea-Based HDAC Inhibitors. <i>ChemMedChem</i> , 2014, 9, 665-670.	3.2	26
122	A novel approach for the discovery of chemically diverse anti-malarial compounds targeting the <i>Plasmodium falciparum</i> Coenzyme A synthesis pathway. <i>Malaria Journal</i> , 2014, 13, 343.	2.3	34
123	Euodenine A: A Small-Molecule Agonist of Human TLR4. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 1252-1275.	6.4	47
124	Lysine Acetylation in Sexual Stage Malaria Parasites Is a Target for Antimalarial Small Molecules. <i>Antimicrobial Agents and Chemotherapy</i> , 2014, 58, 3666-3678.	3.2	62
125	Solving the Supply of Resveratrol Tetramers from Papua New Guinean Rainforest <i>Anisoptera</i> Species That Inhibit Bacterial Type III Secretion Systems. <i>Journal of Natural Products</i> , 2014, 77, 2633-2640.	3.0	16
126	Medicinal Chemistry Optimization of Antiplasmodial Imidazopyridazine Hits from High Throughput Screening of a SoftFocus Kinase Library: Part 1. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 2789-2798.	6.4	43

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127	Facile Synthesis and Preliminary Structure-Activity Analysis of New Sulfonamides Against <i>Trypanosoma brucei</i> . ACS Medicinal Chemistry Letters, 2014, 5, 496-500.	2.8	18
128	2,4-Diaminothienopyrimidines as Orally Active Antimalarial Agents. Journal of Medicinal Chemistry, 2014, 57, 1014-1022.	6.4	34
129	Screening and hit evaluation of a chemical library against blood-stage Plasmodium falciparum. Malaria Journal, 2014, 13, 190.	2.3	47
130	N-Aryl-2-aminobenzimidazoles: Novel, Efficacious, Antimalarial Lead Compounds. Journal of Medicinal Chemistry, 2014, 57, 6642-6652.	6.4	37
131	Pyridyl Benzamides as a Novel Class of Potent Inhibitors for the Kinetoplastid <i>Trypanosoma brucei</i> . Journal of Medicinal Chemistry, 2014, 57, 6393-6402.	6.4	53
132	Repositioning: the fast track to new anti-malarial medicines?. Malaria Journal, 2014, 13, 143.	2.3	36
133	Two-Pronged Attack: Dual Inhibition of Plasmodium falciparum M1 and M17 Metalloaminopeptidases by a Novel Series of Hydroxamic Acid-Based Inhibitors. Journal of Medicinal Chemistry, 2014, 57, 9168-9183.	6.4	52
134	Blood Schizontocidal and Gametocytocidal Activity of 3-Hydroxy- α -arylidene propanehydrazonamides: A New Class of Antiplasmodial Compounds. Journal of Medicinal Chemistry, 2014, 57, 7971-7976.	6.4	13
135	Aminoazabenzimidazoles, a Novel Class of Orally Active Antimalarial Agents. Journal of Medicinal Chemistry, 2014, 57, 5702-5713.	6.4	24
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