

Paul M O'Neill

List of Publications by Year in descending order

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138
papers

8,853
citations

41627

51
h-index

58552

86
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143
all docs

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docs citations

143
times ranked

9736
citing authors

#	ARTICLE	IF	CITATIONS
1	Remdesivir-ivermectin combination displays synergistic interaction with improved in vitro activity against SARS-CoV-2. <i>International Journal of Antimicrobial Agents</i> , 2022, 59, 106542.	1.1	7
2	Dose prediction for repurposing nitazoxanide in SARS-CoV-2 treatment or chemoprophylaxis. <i>British Journal of Clinical Pharmacology</i> , 2021, 87, 2078-2088.	1.1	46
3	Therapeutic Potential of Nitazoxanide: An Appropriate Choice for Repurposing versus SARS-CoV-2?. <i>ACS Infectious Diseases</i> , 2021, 7, 1317-1331.	1.8	37
4	Inhibition mechanism of SARS-CoV-2 main protease by ebiselen and its derivatives. <i>Nature Communications</i> , 2021, 12, 3061.	5.8	149
5	Enantioselective Synthesis and Profiling of Potent, Nonlinear Analogues of Antimalarial Tetraoxanes E209 and N205. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 1077-1085.	1.3	5
6	Anti-Wolbachia drugs for filariasis. <i>Trends in Parasitology</i> , 2021, 37, 1068-1081.	1.5	27
7	Synthesis of Non-symmetrical Dispiro-1,2,4,5-Tetraoxanes and Dispiro-1,2,4-Trioxanes Catalyzed by Silica Sulfuric Acid. <i>Journal of Organic Chemistry</i> , 2021, 86, 10608-10620.	1.7	11
8	Artemisinin inspired synthetic endoperoxide drug candidates: Design, synthesis, and mechanism of action studies. <i>Medicinal Research Reviews</i> , 2021, 41, 3062-3095.	5.0	22
9	Development of Pyrazolopyrimidine Anti-Wolbachia Agents for the Treatment of Filariasis. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 1421-1426.	1.3	5
10	Synthesis, antiviral activity, preliminary pharmacokinetics and structural parameters of thiazolidine amine salts. <i>Future Medicinal Chemistry</i> , 2021, 13, 1731-1741.	1.1	7
11	Machine learning Predicting Ames mutagenicity of small molecules. <i>Journal of Molecular Graphics and Modelling</i> , 2021, 109, 108011.	1.3	11
12	Synthesis, insecticidal activities and resistance in <i>Aedes albopictus</i> and cytotoxicity of novel dihaloacetylated heterocyclic pyrethroids. <i>Pest Management Science</i> , 2020, 76, 636-644.	1.7	15
13	Modification of the cyclopropyl moiety of abacavir provides insight into the structure activity relationship between HLA-B*57:01 binding and T cell activation. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2020, 75, 636-647.	2.7	19
14	Novel Selenium-based compounds with therapeutic potential for SOD1-linked amyotrophic lateral sclerosis. <i>EBioMedicine</i> , 2020, 59, 102980.	2.7	31
15	Prioritization of Anti-SARS-CoV-2 Drug Repurposing Opportunities Based on Plasma and Target Site Concentrations Derived from their Established Human Pharmacokinetics. <i>Clinical Pharmacology and Therapeutics</i> , 2020, 108, 775-790.	2.3	118
16	Ebiselen as template for stabilization of A4V mutant dimer for motor neuron disease therapy. <i>Communications Biology</i> , 2020, 3, 97.	2.0	30
17	Synthesis, insecticidal activity, resistance, photodegradation and toxicity of pyrethroids (A review). <i>Chemosphere</i> , 2020, 254, 126779.	4.2	74
18	Antimalarial Agents as Therapeutic Tools Against Toxoplasmosis—A Short Bridge between Two Distant Illnesses. <i>Molecules</i> , 2020, 25, 1574.	1.7	23

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19	Positively selected modifications in the pore of TbAQP2 allow pentamidine to enter Trypanosoma brucei. ELife, 2020, 9, .	2.8	16
20	Antimalarial activity of primaquine operates via a two-step biochemical relay. Nature Communications, 2019, 10, 3226.	5.8	94
21	Control and regulation of S-Adenosylmethionine biosynthesis by the regulatory $\hat{2}$ subunit and quinolone-based compounds. FEBS Journal, 2019, 286, 2135-2154.	2.2	9
22	Synthesis of MeBmt and related derivatives via syn-selective ATH-DKR. RSC Advances, 2019, 9, 40336-40339.	1.7	7
23	Industrial scale high-throughput screening delivers multiple fast acting macrofilaricides. Nature Communications, 2019, 10, 11.	5.8	93
24	Phosphinic acids: current status and potential for drug discovery. Drug Discovery Today, 2019, 24, 916-929.	3.2	29
25	AWZ1066S, a highly specific anti- <i>Wolbachia</i> drug candidate for a short-course treatment of filariasis. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 1414-1419.	3.3	57
26	Second-generation nitazoxanide derivatives: thiazolidines are effective inhibitors of the influenza A virus. Future Medicinal Chemistry, 2018, 10, 851-862.	1.1	20
27	The cysteine-reactive small molecule ebselen facilitates effective SOD1 maturation. Nature Communications, 2018, 9, 1693.	5.8	71
28	The biological evaluation of fusidic acid and its hydrogenation derivative as antimicrobial and anti-inflammatory agents. Infection and Drug Resistance, 2018, Volume 11, 1945-1957.	1.1	26
29	Potent Antimalarial 2-Pyrazolyl Quinolone Q_{1} Inhibitors with Improved Drug-like Properties. ACS Medicinal Chemistry Letters, 2018, 9, 1205-1210.	1.3	28
30	$\hat{1}$ -Methyl- $\hat{1}$ -phenylsuccinimide ameliorates neurodegeneration in a C. elegans model of TDP-43 proteinopathy. Neurobiology of Disease, 2018, 118, 40-54.	2.1	19
31	X-ray and cryo-EM structures of inhibitor-bound cytochrome bc_{1} complexes for structure-based drug discovery. IUCr, 2018, 5, 200-210.	1.0	23
32	Study of the antimalarial activity of 4-aminoquinoline compounds against chloroquine-sensitive and chloroquine-resistant parasite strains. Journal of Molecular Modeling, 2018, 24, 237.	0.8	24
33	On the ordeal of quinolone preparation via cyclisation of aryl-enamines; synthesis and structure of ethyl 6-methyl-7-iodo-4-(3-iodo-4-methylphenoxy)-quinoline-3-carboxylate. Pure and Applied Chemistry, 2017, 89, 765-780.	0.9	4
34	A tetraoxane-based antimalarial drug candidate that overcomes PfK13-C580Y dependent artemisinin resistance. Nature Communications, 2017, 8, 15159.	5.8	51
35	Rational Design, Synthesis, and Biological Evaluation of Heterocyclic Quinolones Targeting the Respiratory Chain of <i>Mycobacterium tuberculosis</i> . Journal of Medicinal Chemistry, 2017, 60, 3703-3726.	2.9	39
36	Identification and prioritization of novel anti- <i>Wolbachia</i> chemotypes from screening a 10,000-compound diversity library. Science Advances, 2017, 3, eaao1551.	4.7	24

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37	Synthesis and structure-activity relationship of N ⁴ -benzylamine-N ² -isopropyl-quinazoline-2,4-diamines derivatives as potential antibacterial agents. <i>RSC Advances</i> , 2017, 7, 52227-52237.	1.7	12
38	A Click Chemistry-Based Proteomic Approach Reveals that 1,2,4-Trioxolane and Artemisinin Antimalarials Share a Common Protein Alkylation Profile. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 6401-6405.	7.2	76
39	A Click Chemistry-Based Proteomic Approach Reveals that 1,2,4-Trioxolane and Artemisinin Antimalarials Share a Common Protein Alkylation Profile. <i>Angewandte Chemie</i> , 2016, 128, 6511-6515.	1.6	19
40	Molecular Mechanism of Action of Antimalarial Benzoisothiazolones: Species-Selective Inhibitors of the Plasmodium spp. MEP Pathway enzyme, IspD. <i>Scientific Reports</i> , 2016, 6, 36777.	1.6	13
41	Design and Synthesis of Irreversible Analogues of Bardoxolone Methyl for the Identification of Pharmacologically Relevant Targets and Interaction Sites. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 2396-2409.	2.9	37
42	Antimalarial Chemotherapy: Natural Product Inspired Development of Preclinical and Clinical Candidates with Diverse Mechanisms of Action. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 5587-5603.	2.9	59
43	Artemisinin activity-based probes identify multiple molecular targets within the asexual stage of the malaria parasites <i>Plasmodium falciparum</i> 3D7. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, 2080-2085.	3.3	209
44	Small Molecule Inhibitors of Cyclophilin D To Protect Mitochondrial Function as a Potential Treatment for Acute Pancreatitis. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 2596-2611.	2.9	42
45	A Quinoline Carboxamide Antimalarial Drug Candidate Uniquely Targets Plasmodia at Three Stages of the Parasite Life Cycle. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 13504-13506.	7.2	12
46	2-Pyridylquinolone antimalarials with improved antimalarial activity and physicochemical properties. <i>MedChemComm</i> , 2015, 6, 1252-1259.	3.5	14
47	Antimalarial 4(1H)-pyridones bind to the Q site of cytochrome <i>bc₁</i> . <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 755-760.	3.3	90
48	<i>Plasmodium</i> IspD (2-C-Methyl-erythritol 4-Phosphate Cytidyltransferase), an Essential and Druggable Antimalarial Target. <i>ACS Infectious Diseases</i> , 2015, 1, 157-167.	1.8	42
49	From hybrid compounds to targeted drug delivery in antimalarial therapy. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 5120-5130.	1.4	38
50	Integrated transcriptomic and proteomic analyses uncover regulatory roles of Nrf2 in the kidney. <i>Kidney International</i> , 2015, 88, 1261-1273.	2.6	41
51	Carbamoyl Triazoles, Known Serine Protease Inhibitors, Are a Potent New Class of Antimalarials. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 6448-6455.	2.9	17
52	Quinolone-Hydroxyquinoline Tautomerism in Quinolone 3-Esters. Preserving the 4-Oxoquinoline Structure To Retain Antimalarial Activity. <i>Journal of Organic Chemistry</i> , 2015, 80, 12244-12257.	1.7	17
53	Tetraoxane-Pyrimidine Nitrile Hybrids as Dual Stage Antimalarials. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 4916-4923.	2.9	43
54	Rapid kill of malaria parasites by artemisinin and semi-synthetic endoperoxides involves ROS-dependent depolarization of the membrane potential. <i>Journal of Antimicrobial Chemotherapy</i> , 2014, 69, 1005-1016.	1.3	116

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55	Novel Endoperoxide-Based Transmission-Blocking Antimalarials with Liver- and Blood-Schizontocidal Activities. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 108-112.	1.3	40
56	An Endoperoxide-Based Hybrid Approach to Deliver Falcipain Inhibitors Inside Malaria Parasites. <i>ChemMedChem</i> , 2013, 8, 1528-1536.	1.6	32
57	Synthesis and evaluation of the antimalarial, anticancer, and caspase 3 activities of tetraoxane dimers. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 7392-7397.	1.4	19
58	Antimalarial pharmacology and therapeutics of atovaquone. <i>Journal of Antimicrobial Chemotherapy</i> , 2013, 68, 977-985.	1.3	147
59	Artemisinin-Polypyrrole Conjugates: Synthesis, DNA Binding Studies and Preliminary Antiproliferative Evaluation. <i>ChemMedChem</i> , 2013, 8, 709-718.	1.6	7
60	Oxidative Bioactivation of Abacavir in Subcellular Fractions of Human Antigen Presenting Cells. <i>Chemical Research in Toxicology</i> , 2013, 26, 1064-1072.	1.7	12
61	Pyrethroid activity-based probes for profiling cytochrome P450 activities associated with insecticide interactions. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 19766-19771.	3.3	33
62	Targeting the mitochondrial electron transport chain of <i>Plasmodium falciparum</i> : new strategies towards the development of improved antimalarials for the elimination era. <i>Future Medicinal Chemistry</i> , 2013, 5, 1573-1591.	1.1	55
63	Cytochrome b Mutation Y268S Conferring Atovaquone Resistance Phenotype in Malaria Parasite Results in Reduced Parasite bc1 Catalytic Turnover and Protein Expression. <i>Journal of Biological Chemistry</i> , 2012, 287, 9731-9741.	1.6	77
64	HDQ, a Potent Inhibitor of <i>Plasmodium falciparum</i> Proliferation, Binds to the Quinone Reduction Site of the Cytochrome bc 1 Complex. <i>Antimicrobial Agents and Chemotherapy</i> , 2012, 56, 3739-3747.	1.4	53
65	Identification, Design and Biological Evaluation of Bisaryl Quinolones Targeting <i>Plasmodium falciparum</i> Type II NADH:Quinone Oxidoreductase (PfNDH2). <i>Journal of Medicinal Chemistry</i> , 2012, 55, 1831-1843.	2.9	94
66	Identification, Design and Biological Evaluation of Heterocyclic Quinolones Targeting <i>Plasmodium falciparum</i> Type II NADH:Quinone Oxidoreductase (PfNDH2). <i>Journal of Medicinal Chemistry</i> , 2012, 55, 1844-1857.	2.9	51
67	Identification of Novel Antimalarial Chemotypes via Chemoinformatic Compound Selection Methods for a High-Throughput Screening Program against the Novel Malarial Target, PfNDH2: Increasing Hit Rate via Virtual Screening Methods. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 3144-3154.	2.9	23
68	Examination of the Cytotoxic and Embryotoxic Potential and Underlying Mechanisms of Next-Generation Synthetic Trioxolane and Tetraoxane Antimalarials. <i>Molecular Medicine</i> , 2012, 18, 1045-1055.	1.9	12
69	The MEP pathway and the development of inhibitors as potential anti-infective agents. <i>MedChemComm</i> , 2012, 3, 418.	3.5	41
70	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.	3.3	143
71	Convenient Syntheses of Benzo-Fluorinated Dibenz[<i>b</i>], [<i>f</i>]azepines: Rearrangements of Isatins, Acridines, and Indoles. <i>Organic Letters</i> , 2011, 13, 5592-5595.	2.4	30
72	Comparison of the Reactivity of Antimalarial 1,2,4,5-Tetraoxanes with 1,2,4-Trioxolanes in the Presence of Ferrous Iron Salts, Heme, and Ferrous Iron Salts/Phosphatidylcholine. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 6443-6455.	2.9	47

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73	Second generation analogues of RKA182: synthetic tetraoxanes with outstanding in vitro and in vivo antimalarial activities. <i>MedChemComm</i> , 2011, 2, 661.	3.5	28
74	Cytochrome P450 6M2 from the malaria vector <i>Anopheles gambiae</i> metabolizes pyrethroids: Sequential metabolism of deltamethrin revealed. <i>Insect Biochemistry and Molecular Biology</i> , 2011, 41, 492-502.	1.2	217
75	Antimalarial Mannoxanes: Hybrid Antimalarial Drugs with Outstanding Oral Activity Profiles and A Potential Dual Mechanism of Action. <i>ChemMedChem</i> , 2011, 6, 1357-1361.	1.6	25
76	Synthesis and Antimalarial Activities of a Diverse Set of Triazole-Containing Furamide Analogues. <i>ChemMedChem</i> , 2011, 6, 2094-2108.	1.6	26
77	The Role of Heme and the Mitochondrion in the Chemical and Molecular Mechanisms of Mammalian Cell Death Induced by the Artemisinin Antimalarials. <i>Journal of Biological Chemistry</i> , 2011, 286, 987-996.	1.6	137
78	A novel drug for uncomplicated malaria: Targeted high throughput screening (HTS) against the type II NADH:ubiquinone oxidoreductase (PfNDH2) of <i>Plasmodium falciparum</i> . <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2010, 1797, 80.	0.5	0
79	Identification of a 1,2,4,5-Tetraoxane Antimalarial Drug-Development Candidate (RKA182) with Superior Properties to the Semisynthetic Artemisinins. <i>Angewandte Chemie - International Edition</i> , 2010, 49, 5693-5697.	7.2	111
80	Inhibiting <i>Plasmodium</i> cytochrome bc1: a complex issue. <i>Current Opinion in Chemical Biology</i> , 2010, 14, 440-446.	2.8	97
81	Design, synthesis and antimalarial/anticancer evaluation of spermidine linked artemisinin conjugates designed to exploit polyamine transporters in <i>Plasmodium falciparum</i> and HL-60 cancer cell lines. <i>Bioorganic and Medicinal Chemistry</i> , 2010, 18, 2586-2597.	1.4	51
82	Modular Synthesis and in Vitro and in Vivo Antimalarial Assessment of C-10 Pyrrole Mannich Base Derivatives of Artemisinin. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 633-640.	2.9	52
83	The Molecular Mechanism of Action of Artemisinin- The Debate Continues. <i>Molecules</i> , 2010, 15, 1705-1721.	1.7	474
84	Endoperoxide Carbonyl Falcipain 2/3 Inhibitor Hybrids: Toward Combination Chemotherapy of Malaria through a Single Chemical Entity. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 8202-8206.	2.9	35
85	Rationale Design of Biotinylated Antimalarial Endoperoxide Carbon Centered Radical Prodrugs for Applications in Proteomics. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 4555-4559.	2.9	29
86	A novel drug for uncomplicated malaria: targeted high throughput screening (HTS) against the type II NADH:ubiquinone oxidoreductase (PfNdh2) of <i>Plasmodium falciparum</i> . <i>Malaria Journal</i> , 2010, 9, .	0.8	2
87	Comparative preclinical drug metabolism and pharmacokinetic evaluation of novel 4-aminoquinoline anti-malarials. <i>Journal of Pharmaceutical Sciences</i> , 2009, 98, 362-377.	1.6	16
88	Synthesis and biological evaluation of extraordinarily potent C-10 carba artemisinin dimers against <i>P. falciparum</i> malaria parasites and HL-60 cancer cells. <i>Bioorganic and Medicinal Chemistry</i> , 2009, 17, 1325-1338.	1.4	58
89	Semi-synthetic and synthetic 1,2,4-trioxaquinones and 1,2,4-trioxolaquinones: synthesis, preliminary SAR and comparison with acridine endoperoxide conjugates. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009, 19, 2038-2043.	1.0	64
90	Antitumour and antimalarial activity of artemisinin-acridine hybrids. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009, 19, 2033-2037.	1.0	50

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91	Candidate Selection and Preclinical Evaluation of <i>N</i> - <i>tert</i> -Butyl Isoquine (GSK369796), An Affordable and Effective 4-Aminoquinoline Antimalarial for the 21st Century. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 1408-1415.	2.9	80
92	Synthesis, Antimalarial Activity, and Preclinical Pharmacology of a Novel Series of 4-Fluoro and 4-Chloro Analogues of Amodiaquine. Identification of a Suitable "Back-Up" Compound for <i>N</i> - <i>tert</i> -Butyl Isoquine. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 1828-1844.	2.9	56
93	An efficient route into synthetically challenging bridged achiral 1,2,4,5-tetraoxanes with antimalarial activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 1720-1724.	1.0	30
94	Piperidine dispiro-1,2,4-trioxane analogues. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 5804-5808.	1.0	27
95	Two-Step Synthesis of Achiral Dispiro-1,2,4,5-tetraoxanes with Outstanding Antimalarial Activity, Low Toxicity, and High-Stability Profiles. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 2170-2177.	2.9	78
96	Acridinediones: Selective and Potent Inhibitors of the Malaria Parasite Mitochondrial bc1 Complex. <i>Molecular Pharmacology</i> , 2008, 73, 1347-1355.	1.0	85
97	Evidence for the Involvement of Carbon-centered Radicals in the Induction of Apoptotic Cell Death by Artemisinin Compounds. <i>Journal of Biological Chemistry</i> , 2007, 282, 9372-9382.	1.6	164
98	Evidence for a Common Non-Heme Chelatable-Dependent Activation Mechanism for Semisynthetic and Synthetic Endoperoxide Antimalarial Drugs. <i>Angewandte Chemie - International Edition</i> , 2007, 46, 6278-6283.	7.2	116
99	Anticancer activity of artemisinin-derived trioxanes. <i>Expert Opinion on Therapeutic Patents</i> , 2006, 16, 1665-1672.	2.4	41
100	Design and synthesis of orally active dispiro 1,2,4,5-tetraoxanes; synthetic antimalarials with superior activity to artemisinin. <i>Organic and Biomolecular Chemistry</i> , 2006, 4, 4431.	1.5	83
101	Diels-Alder/thiol-olefin co-oxygenation approach to antimalarials incorporating the 2,3-dioxabicyclo[3.3.1]nonane pharmacophore. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006, 16, 2991-2995.	1.0	19
102	Synthesis of 1,2,4-trioxepanes via application of thiol-olefin Co-oxygenation methodology. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006, 16, 6124-6130.	1.0	13
103	Functional Characterization and Target Validation of Alternative Complex I of <i>Plasmodium falciparum</i> Mitochondria. <i>Antimicrobial Agents and Chemotherapy</i> , 2006, 50, 1841-1851.	1.4	120
104	A Medicinal Chemistry Perspective on 4-Aminoquinoline Antimalarial Drugs. <i>Current Topics in Medicinal Chemistry</i> , 2006, 6, 479-507.	1.0	104
105	Enantiomeric 1,2,4-Trioxanes Display Equivalent <i>in vitro</i> Antimalarial Activity Versus <i>Plasmodium falciparum</i> Malaria Parasites: Implications for the Molecular Mechanism of Action of the Artemisinins. <i>ChemBioChem</i> , 2005, 6, 2048-2054.	1.3	49
106	The therapeutic potential of semi-synthetic artemisinin and synthetic endoperoxide antimalarial agents. <i>Expert Opinion on Investigational Drugs</i> , 2005, 14, 1117-1128.	1.9	37
107	Current drug development portfolio for antimalarial therapies. <i>Current Opinion in Pharmacology</i> , 2005, 5, 473-478.	1.7	46
108	A worthy adversary for malaria. <i>Nature</i> , 2004, 430, 838-839.	13.7	49

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109	Design and Synthesis of Endoperoxide Antimalarial Prodrug Models. <i>Angewandte Chemie - International Edition</i> , 2004, 43, 4193-4197.	7.2	56
110	A Medicinal Chemistry Perspective on Artemisinin and Related Endoperoxides. <i>ChemInform</i> , 2004, 35, no.	0.1	0
111	Knowledge of the Proposed Chemical Mechanism of Action and Cytochrome P450 Metabolism of Antimalarial Trioxanes Like Artemisinin Allows Rational Design of New Antimalarial Peroxides. <i>ChemInform</i> , 2004, 35, no.	0.1	0
112	Antimalarial and Antitumor Evaluation of Novel C-10 Non-Acetal Dimers of 10 ^β -(2-Hydroxyethyl)deoxoartemisinin. <i>Journal of Medicinal Chemistry</i> , 2004, 47, 1290-1298.	2.9	97
113	Application of Thiol ^α Olefin Co-oxygenation Methodology to a New Synthesis of the 1,2,4-Trioxane Pharmacophore. <i>Organic Letters</i> , 2004, 6, 3035-3038.	2.4	58
114	A Medicinal Chemistry Perspective on Artemisinin and Related Endoperoxides. <i>Journal of Medicinal Chemistry</i> , 2004, 47, 2945-2964.	2.9	505
115	Knowledge of the Proposed Chemical Mechanism of Action and Cytochrome P450 Metabolism of Antimalarial Trioxanes Like Artemisinin Allows Rational Design of New Antimalarial Peroxides. <i>Accounts of Chemical Research</i> , 2004, 37, 397-404.	7.6	214
116	Antimalarial chemotherapy: young guns or back to the future?. <i>Trends in Parasitology</i> , 2003, 19, 479-487.	1.5	79
117	Co(thd) ₂ : a superior catalyst for aerobic epoxidation and hydroperoxysilylation of unactivated alkenes: application to the synthesis of spiro-1,2,4-trioxanes. <i>Tetrahedron Letters</i> , 2003, 44, 8135-8138.	0.7	69
118	Isoquine and Related Amodiaquine Analogues: A New Generation of Improved 4-Aminoquinoline Antimalarials. <i>Journal of Medicinal Chemistry</i> , 2003, 46, 4933-4945.	2.9	130
119	Mechanism-Based Design of Parasite-Targeted Artemisinin Derivatives: Synthesis and Antimalarial Activity of New Diamine Containing Analogues. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 1052-1063.	2.9	116
120	Novel Short Chain Chloroquine Analogues Retain Activity Against Chloroquine Resistant K1 Plasmodium falciparum. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 4975-4983.	2.9	121
121	Synthesis, Antimalarial Activity, Biomimetic Iron(II) Chemistry, and in Vivo Metabolism of Novel, Potent C-10-Phenoxy Derivatives of Dihydroartemisinin. <i>Journal of Medicinal Chemistry</i> , 2001, 44, 58-68.	2.9	92
122	METABOLISM OFFLUORINE-CONTAININGDRUGS. <i>Annual Review of Pharmacology and Toxicology</i> , 2001, 41, 443-470.	4.2	550
123	Regioselective Mukaiyama hydroperoxysilylation of 2-alkyl- or 2-aryl-prop-2-en-1-ols: application to a new synthesis of 1,2,4-trioxanes. <i>Tetrahedron Letters</i> , 2001, 42, 4569-4571.	0.7	54
124	Biomimetic Fe(II)-Mediated Degradation of Arteflene (Ro-42-1611). The First EPR Spin-Trapping Evidence for the Previously Postulated Secondary Carbon-Centered Cyclohexyl Radical. <i>Journal of Organic Chemistry</i> , 2000, 65, 1578-1582.	1.7	59
125	Asymmetric syntheses of enantiomeric 3-p-fluorophenyl 1,2,4-trioxane analogues of the antimalarial artemisinin. <i>Tetrahedron Letters</i> , 1999, 40, 9133-9136.	0.7	22
126	Novel, Potent, Semisynthetic Antimalarial Carba Analogues of the First-Generation 1,2,4-Trioxane Artemether. <i>Journal of Medicinal Chemistry</i> , 1999, 42, 5487-5493.	2.9	58

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127	New 4-Aminoquinoline Mannich Base Antimalarials. 1. Effect of an Alkyl Substituent in the 5-Position of the 4-Hydroxyanilino Side Chain. <i>Journal of Medicinal Chemistry</i> , 1999, 42, 2747-2751.	2.9	58
128	4-Aminoquinolines—Past, present, and future; A chemical perspective. , 1998, 77, 29-58.		242
129	Synthesis of the 8-aminoquinoline antimalarial 5-fluoroprimaquine. <i>Tetrahedron</i> , 1998, 54, 4615-4622.	1.0	30
130	Safety assessment of peroxide antimalarials: clinical and chemical perspectives. <i>British Journal of Clinical Pharmacology</i> , 1998, 46, 521-529.	1.1	41
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138	Unprecedented Convergent Synthesis of Sugar-Functionalization of Phosphinic Acids under Metal-Free Conditions. <i>ACS Omega</i> , 0, , .	1.6	4