## Scott G Franzblau

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6557742/publications.pdf

Version: 2024-02-01

326 papers 15,275 citations

64 h-index 99 g-index

348 all docs 348 docs citations

times ranked

348

15486 citing authors

#	Article	IF	CITATIONS
1	Rapid, Low-Technology MIC Determination with Clinical <i>Mycobacterium tuberculosis</i> Isolates by Using the Microplate Alamar Blue Assay. Journal of Clinical Microbiology, 1998, 36, 362-366.	3.9	810
2	Low-Oxygen-Recovery Assay for High-Throughput Screening of Compounds against Nonreplicating Mycobacterium tuberculosis. Antimicrobial Agents and Chemotherapy, 2007, 51, 1380-1385.	3.2	286
3	Structureâ^'Activity Relationships for a Series of Quinoline-Based Compounds Active against Replicating and Nonreplicating <i>Mycobacterium tuberculosis</i> . Journal of Medicinal Chemistry, 2009, 52, 2109-2118.	6.4	275
4	Thiosemicarbazones, semicarbazones, dithiocarbazates and hydrazide/hydrazones: Anti – Mycobacterium tuberculosis activity and cytotoxicity. European Journal of Medicinal Chemistry, 2010, 45, 1898-1905.	5.5	272
5	Improved Green Fluorescent Protein Reporter Gene-BasedMicroplate Screening for Antituberculosis Compounds by Utilizing anAcetamidasePromoter. Antimicrobial Agents and Chemotherapy, 2003, 47, 3682-3687.	3.2	241
6	Antimycobacterial Plant Terpenoids. Planta Medica, 2001, 67, 685-694.	1.3	212
7	Targeting mycobacterium protein tyrosine phosphatase B for antituberculosis agents. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 4573-4578.	7.1	211
8	Comprehensive analysis of methods used for the evaluation of compounds against Mycobacterium tuberculosis. Tuberculosis, 2012, 92, 453-488.	1.9	193
9	In Vitro and In Vivo Activities of Macrolide Derivatives against <i>Mycobacterium tuberculosis</i> Antimicrobial Agents and Chemotherapy, 2005, 49, 1447-1454.	3.2	191
10	Identification of a small molecule with activity against drug-resistant and persistent tuberculosis. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, E2510-7.	7.1	188
11	Novel Insights into the Mechanism of Inhibition of MmpL3, a Target of Multiple Pharmacophores in Mycobacterium tuberculosis. Antimicrobial Agents and Chemotherapy, 2014, 58, 6413-6423.	3.2	174
12	Advent of Imidazo[1,2- <i>a</i> ]pyridine-3-carboxamides with Potent Multi- and Extended Drug Resistant Antituberculosis Activity. ACS Medicinal Chemistry Letters, 2011, 2, 466-470.	2.8	161
13	Drug Targeting Mycobacterium tuberculosis Cell Wall Synthesis: Genetics of dTDP-Rhamnose Synthetic Enzymes and Development of a Microtiter Plate-Based Screen for Inhibitors of Conversion of dTDP-Glucose to dTDP-Rhamnose. Antimicrobial Agents and Chemotherapy, 2001, 45, 1407-1416.	3.2	151
14	The Cyclic Peptide Ecumicin Targeting ClpC1 Is Active against Mycobacterium tuberculosis In Vivo. Antimicrobial Agents and Chemotherapy, 2015, 59, 880-889.	3.2	148
15	Synthesis, antimalarial and antitubercular activity of acetylenic chalcones. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 942-944.	2.2	146
16	Diguanidino and "Reversed―Diamidino 2,5-Diarylfurans as Antimicrobial Agents. Journal of Medicinal Chemistry, 2001, 44, 1741-1748.	6.4	135
17	Manzamine B and E and Ircinal A Related Alkaloids from an IndonesianAcanthostrongylophoraSponge and Their Activity against Infectious, Tropical Parasitic, and Alzheimer's Diseases. Journal of Natural Products, 2006, 69, 1034-1040.	3.0	129
18	Antitubercular constituents from the hexane fraction of Morinda citrifolia Linn. (Rubiaceae). Phytotherapy Research, 2002, 16, 683-685.	5.8	124

#	Article	IF	Citations
19	New perspectives on natural products in TB drug research. Life Sciences, 2005, 78, 485-494.	4.3	120
20	Antimycobacterial Activity of (E)-Phytol and Derivatives: A Preliminary Structure-Activity Study. Planta Medica, 1998, 64, 2-4.	1.3	118
21	Synthesis and Activity of Carbazole Derivatives AgainstMycobacterium tuberculosis. ChemMedChem, 2006, 1, 812-815.	3.2	108
22	Discovery of Selective Menaquinone Biosynthesis Inhibitors against Mycobacterium tuberculosis. Journal of Medicinal Chemistry, 2012, 55, 3739-3755.	6.4	106
23	Synthesis and Structurea "Activity Studies of Biphenyl Analogues of the Tuberculosis Drug $(6 < i > S <  i>)-2-Nitro-6-{[4-(trifluoromethoxy)benzyl]oxy}-6,7-dihydro-5 < i>H <  i>-imidazo[2,1-b <  i>)[1,3]oxazine (PA-824). Journal of Medicinal Chemistry, 2010, 53, 282-294.$	6.4	104
24	5- <i>tert</i> -Butyl- <i>N</i> -pyrazol-4-yl-4,5,6,7-tetrahydrobenzo[ <i>d</i> ]isoxazole-3-carboxamide Derivatives as Novel Potent Inhibitors of <i>Mycobacterium tuberculosis</i> Synthetase: Initiating a Quest for New Antitubercular Drugs. Journal of Medicinal Chemistry, 2008, 51, 1999-2002.	6.4	102
25	Structure–activity relationship of new anti-tuberculosis agents derived from oxazoline and oxazole benzyl esters. European Journal of Medicinal Chemistry, 2010, 45, 1703-1716.	5.5	99
26	Design, Synthesis, and SAR Studies of Mefloquine-Based Ligands as Potential Antituberculosis Agents. ChemMedChem, 2006, 1, 593-597.	3.2	98
27	Design, Synthesis, and Structure–Activity Relationship Studies of Tryptanthrins As Antitubercular Agents. Journal of Natural Products, 2013, 76, 354-367.	3.0	98
28	New tuberculosis drug targets, their inhibitors, and potential therapeutic impact. Translational Research, 2020, 220, 68-97.	5.0	97
29	Antimycobacterial natural products: synthesis and preliminary biological evaluation of the oxazole-containing alkaloid texaline. Tetrahedron Letters, 2005, 46, 7355-7357.	1.4	96
30	Generation and exploration of new classes of antitubercular agents: The optimization of oxazolines, oxazoles, thiazolines, thiazoles to imidazo[1,2-a]pyridines and isomeric 5,6-fused scaffolds. Bioorganic and Medicinal Chemistry, 2012, 20, 2214-2220.	3.0	96
31	QSAR-driven design, synthesis and discovery of potent chalcone derivatives with antitubercular activity. European Journal of Medicinal Chemistry, 2017, 137, 126-138.	5.5	96
32	Synthesis and evaluation of anti-tubercular and antibacterial activities of new 4-(2,6-dichlorobenzyloxy)phenyl thiazole, oxazole and imidazole derivatives. Part 2. European Journal of Medicinal Chemistry, 2012, 49, 164-171.	5.5	95
33	Antimycobacterial Eudesmanolides from Inula helenium and Rudbeckia subtomentosa. Planta Medica, 1999, 65, 351-355.	1.3	94
34	Manadomanzamines A and B:Â A Novel Alkaloid Ring System with Potent Activity against Mycobacteria and HIV-1. Journal of the American Chemical Society, 2003, 125, 13382-13386.	13.7	94
35	A microbiological assessment of novel nitrofuranylamides as anti-tuberculosis agents. Journal of Antimicrobial Chemotherapy, 2008, 62, 1037-1045.	3.0	94
36	Antimycobacterial Cycloartanes fromBorrichiafrutescens. Journal of Natural Products, 1996, 59, 1131-1136.	3.0	92

#	Article	IF	CITATIONS
37	From Serendipity to Rational Antituberculosis Drug Discovery of Mefloquine-Isoxazole Carboxylic Acid Esters. Journal of Medicinal Chemistry, 2009, 52, 6966-6978.	6.4	92
38	Hapalindole-related alkaloids from the cultured cyanobacterium Fischerella ambigua. Phytochemistry, 2010, 71, 2116-2123.	2.9	90
39	Improved BM212 MmpL3 Inhibitor Analogue Shows Efficacy in Acute Murine Model of Tuberculosis Infection. PLoS ONE, 2013, 8, e56980.	2.5	90
40	Transition metals in organic synthesis - Part 83#: Synthesis and pharmacological potential of carbazoles. Medicinal Chemistry Research, 2008, 17, 374-385.	2.4	89
41	Utilization of microbial iron assimilation processes for the development of new antibiotics and inspiration for the design of new anticancer agents. BioMetals, 2009, 22, 61-75.	4.1	89
42	Synthesis, Reduction Potentials, and Antitubercular Activity of Ring A/B Analogues of the Bioreductive Drug (6 <i>&gt;S</i> )-2-Nitro-6-{[4-(trifluoromethoxy)benzyl]oxy}-6,7-dihydro-5 <i>H</i> -imidazo[2,1- <i>b</i> ][1,3]oxazine (PA-824). Journal of Medicinal Chemistry, 2009, 52, 637-645.	6.4	88
43	Discovery of Novel Oral Protein Synthesis Inhibitors of Mycobacterium tuberculosis That Target Leucyl-tRNA Synthetase. Antimicrobial Agents and Chemotherapy, 2016, 60, 6271-6280.	3.2	88
44	Evaluation of gyrase B as a drug target in Mycobacterium tuberculosis. Journal of Antimicrobial Chemotherapy, 2012, 67, 415-421.	3.0	87
45	Antitubercular Constituents of Valeriana laxiflora. Planta Medica, 2004, 70, 509-514.	1.3	85
46	Antimycobacterial evaluation of germacranolides in honour of professor G.H. Neil Towers 75th birthday. Phytochemistry, 1998, 49, 559-564.	2.9	84
47	Microplate Alamar Blue Assay (MABA) and Low Oxygen Recovery Assay (LORA) for Mycobacterium tuberculosis. Methods in Molecular Biology, 2015, 1285, 281-292.	0.9	84
48	Synthesis and Evaluation of Nitrofuranylamides as Novel Antituberculosis Agents. Journal of Medicinal Chemistry, 2004, 47, 5276-5283.	6.4	81
49	Antimycobacterial terpenoids from Juniperus communis L. (Cuppressaceae). Journal of Ethnopharmacology, 2009, 126, 500-505.	4.1	81
50	Synthesis and Structureâ^'activity Relationships of Antitubercular 2-Nitroimidazooxazines Bearing Heterocyclic Side Chains. Journal of Medicinal Chemistry, 2010, 53, 855-866.	6.4	81
51	Anti-Tuberculosis Constituents from the Stem Bark of Micromelum hirsutum. Planta Medica, 2005, 71, 261-267.	1.3	80
52	Synthesis and Structureâ^'Activity Relationships of Aza- and Diazabiphenyl Analogues of the Antitubercular Drug (6 <i>S</i> )-2-Nitro-6-{[4-(trifluoromethoxy)benzyl]oxy}-6,7-dihydro-5 <i>H</i> -imidazo[2,1- <i>b</i> ][1,3]oxazine (PA-824). Journal of Medicinal Chemistry, 2010, 53, 8421-8439.	6.4	80
53	Ethnobotany/ethnopharmacology and mass bioprospecting: Issues on intellectual property and benefit-sharing. Journal of Ethnopharmacology, 2005, 100, 15-22.	4.1	79
54	Identification of Novel Inhibitors of Nonreplicating Mycobacterium tuberculosis Using a Carbon Starvation Model. ACS Chemical Biology, 2013, 8, 2224-2234.	3.4	79

#	Article	IF	CITATIONS
55	Antimycobacterial Activity of Substituted Isosteres of Pyridine- and Pyrazinecarboxylic Acids. Journal of Medicinal Chemistry, 1998, 41, 2436-2438.	6.4	77
56	Antimycobacterial Compounds fromPipersanctumâ€. Journal of Natural Products, 2004, 67, 1961-1968.	3.0	77
57	Antimalarial Bromophycolides Jâ^'Q from the Fijian Red Alga <i>Callophycus serratus</i> . Journal of Organic Chemistry, 2009, 74, 2736-2742.	3.2	77
58	Design, Synthesis, and Pharmacological Evaluation of Mefloquineâ€Based Ligands as Novel Antituberculosis Agents. ChemMedChem, 2007, 2, 1624-1630.	3.2	73
59	Indole alkaloids from the leaves of Philippine Alstonia scholaris. Phytochemistry, 2005, 66, 1158-1162.	2.9	72
60	Antimycobacterial Ergosterol-5,8-endoperoxide from Ajuga remota. Planta Medica, 1999, 65, 732-734.	1.3	70
61	Lahorenoic Acids A–C, <i>ortho</i> -Dialkyl-Substituted Aromatic Acids from the Biocontrol Strain <i>Pseudomonas aurantiaca</i> PB-St2. Journal of Natural Products, 2013, 76, 135-141.	3.0	70
62	Effective Treatment of Acute and Chronic Murine Tuberculosis with Liposome-Encapsulated Clofazimine. Antimicrobial Agents and Chemotherapy, 1999, 43, 1638-1643.	3.2	69
63	3,5-Dialkoxypyridine analogues of bedaquiline are potent antituberculosis agents with minimal inhibition of the hERG channel. Bioorganic and Medicinal Chemistry, 2019, 27, 1292-1307.	3.0	69
64	Antimycobacterial Activity of Substituted Isosteres of Pyridine- and Pyrazinecarboxylic Acids. 2.1. Journal of Medicinal Chemistry, 2001, 44, 1560-1563.	6.4	68
65	Ruthenium(II) phosphine/diimine/picolinate complexes: Inorganic compounds as agents against tuberculosis. European Journal of Medicinal Chemistry, 2011, 46, 5099-5107.	5.5	68
66	Rufomycin Targets ClpC1 Proteolysis in Mycobacterium tuberculosis and M. abscessus. Antimicrobial Agents and Chemotherapy, 2019, 63, .	3.2	68
67	Bioactive metabolites of Diaporthe sp. P133, an endophytic fungus isolated from Pandanus amaryllifolius. Journal of Natural Medicines, 2011, 65, 606-609.	2.3	67
68	Synthesis and Structure–Activity Relationships of Varied Ether Linker Analogues of the Antitubercular Drug (6 <i>&gt;S</i> )-2-Nitro-6-{[4-(trifluoromethoxy)benzyl]oxy}-6,7-dihydro-5 <i>H</i> -imidazo[2,1- <i>b</i> ][1,3]oxazine (PA-824). Journal of Medicinal Chemistry, 2011, 54, 6563-6585.	6.4	66
69	Facile transformation of Biginelli pyrimidin-2(1H)-ones to pyrimidines. In vitro evaluation as inhibitors of Mycobacterium tuberculosis and modulators of cytostatic activity. European Journal of Medicinal Chemistry, 2011, 46, 2290-2294.	5.5	66
70	6-Cyano Analogues of Bedaquiline as Less Lipophilic and Potentially Safer Diarylquinolines for Tuberculosis. ACS Medicinal Chemistry Letters, 2017, 8, 1019-1024.	2.8	66
71	Unbiased evaluation of bioactive secondary metabolites in complex matrices. Fìtoterapìâ, 2012, 83, 1218-1225.	2.2	65
72	Arrival of Imidazo[2,1- <i>b</i> ]thiazole-5-carboxamides: Potent Anti-tuberculosis Agents That Target QcrB. ACS Infectious Diseases, 2016, 2, 393-398.	3.8	64

#	Article	IF	CITATIONS
73	Structure-activity relationships for analogs of the tuberculosis drug bedaquiline with the naphthalene unit replaced by bicyclic heterocycles. Bioorganic and Medicinal Chemistry, 2018, 26, 1797-1809.	3.0	63
74	Assessment of Antimycobacterial Activity of a Series of Mainly Marine Derived Natural Products. Planta Medica, 2000, 66, 337-342.	1.3	62
75	12,34-Oxamanzamines, novel biocatalytic and natural products from manzamine producing Indo-Pacific sponges. Tetrahedron, 2002, 58, 7397-7402.	1.9	62
76	Anti-infective Discorhabdins from a Deep-Water Alaskan Sponge of the Genus <i>Latrunculia</i> Journal of Natural Products, 2010, 73, 383-387.	3.0	61
77	Synthesis and antituberculosis activity of novel mefloquine-isoxazole carboxylic esters as prodrugs. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 1263-1268.	2.2	60
78	Role of antibiotic ligand in nascent peptide-dependent ribosome stalling. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 10496-10501.	7.1	60
79	Purityâ^'Activity Relationships of Natural Products: The Case of Anti-TB Active Ursolic Acid. Journal of Natural Products, 2008, 71, 1742-1748.	3.0	59
80	In vivo and in vitro evaluation of highly specific thiolate carrier group copper(II) and zinc(II) complexes on Ehrlich ascites carcinoma tumor model. European Journal of Medicinal Chemistry, 2010, 45, 5438-5451.	5 <b>.</b> 5	59
81	Antitubercular Activity of Triterpenoids from Asteraceae Flowers. Biological and Pharmaceutical Bulletin, 2005, 28, 158-160.	1.4	58
82	Design, synthesis and investigation on the structure–activity relationships of N-substituted 2-aminothiazole derivatives as antitubercular agents. European Journal of Medicinal Chemistry, 2014, 72, 26-34.	5 <b>.</b> 5	58
83	An Antimicrobial Guanidine-Bearing Sesterterpene from the Cultured Cyanobacterium Scytonema sp Journal of Natural Products, 2009, 72, 2043-2045.	3.0	57
84	Searching for New Cures for Tuberculosis: Design, Synthesis, and Biological Evaluation of 2-Methylbenzothiazoles. Journal of Medicinal Chemistry, 2009, 52, 6757-6767.	6.4	57
85	Rational Design of 5-Phenyl-3-isoxazolecarboxylic Acid Ethyl Esters as Growth Inhibitors of <i>Mycobacterium tuberculosis</i> . A Potent and Selective Series for Further Drug Development. Journal of Medicinal Chemistry, 2010, 53, 678-688.	6.4	57
86	The Oxidation-sensing Regulator (MosR) Is a New Redox-dependent Transcription Factor in Mycobacterium tuberculosis. Journal of Biological Chemistry, 2012, 287, 37703-37712.	3.4	57
87	A New Antitubercular Mulinane Diterpenoid from Azorella madreporica Clos. Journal of Natural Products, 1998, 61, 965-968.	3.0	56
88	Counter-current chromatography based analysis of synergy in an anti-tuberculosis ethnobotanical. Journal of Chromatography A, 2007, 1151, 211-215.	3.7	56
89	In vitro and in vivo antimycobacterial activities of ketone and amide derivatives of quinoxaline 1,4-di-N-oxide. Journal of Antimicrobial Chemotherapy, 2008, 62, 547-554.	3.0	55
90	Discovery of a capuramycin analog that kills nonreplicating Mycobacterium tuberculosis and its synergistic effects with translocase I inhibitors. Journal of Antibiotics, 2015, 68, 271-278.	2.0	55

#	Article	IF	Citations
91	Syntheses and evaluation of benzodiazaborine compounds against M. tuberculosis H37Rv in vitro. Bioorganic and Medicinal Chemistry Letters, 1998, 8, 843-846.	2.2	54
92	Activity of 7-methyljuglone in combination with antituberculous drugs against Mycobacterium tuberculosis. Phytomedicine, 2006, 13, 630-635.	5.3	54
93	Design, Syntheses, and Anti-TB Activity of 1,3-Benzothiazinone Azide and Click Chemistry Products Inspired by BTZ043. ACS Medicinal Chemistry Letters, 2016, 7, 266-270.	2.8	54
94	Evaluation of antiprotozoal and antimycobacterial activities of the resin glycosides and the other metabolites of Scrophularia cryptophila. Phytomedicine, 2008, 15, 209-215.	<b>5.</b> 3	53
95	Preparation of aminoglycoside-loaded chitosan nanoparticles using dextran sulphate as a counterion. Journal of Microencapsulation, 2009, 26, 346-354.	2.8	53
96	Bioactive Bromophycolides Râ^'U from the Fijian Red Alga <i>Callophycus serratus</i> . Journal of Natural Products, 2010, 73, 275-278.	3.0	53
97	Structure–Activity Relationships for Amide-, Carbamate-, And Urea-Linked Analogues of the Tuberculosis Drug (6 < i > S <   i > )-2-Nitro-6-{[4-(trifluoromethoxy)benzyl]oxy}-6,7-dihydro-5 < i > H <   i > -imidazo[2,1- <i> b &lt;   i &gt; )[1,3]oxazine (PA-824). Journal of Medicinal Chemistry, 2012, 55, 312-326.</i>	6.4	53
98	Phomapyrrolidones A–C, Antitubercular Alkaloids from the Endophytic Fungus <i>Phoma</i> sp. NRRL 46751. Journal of Natural Products, 2013, 76, 1860-1865.	3.0	53
99	ICAT-based comparative proteomic analysis of non-replicating persistent Mycobacterium tuberculosis. Tuberculosis, 2006, 86, 445-460.	1.9	52
100	Callophycoic Acids and Callophycols from the Fijian Red Alga <i>Callophycus serratus</i> . Journal of Organic Chemistry, 2007, 72, 7343-7351.	3.2	52
101	Agelasine F from a Philippine Agelas sp. Sponge Exhibits in vitro Antituberculosis Activity. Planta Medica, 2000, 66, 364-365.	1.3	51
102	Enhancing Hit Identification in Mycobacterium tuberculosis Drug Discovery Using Validated Dual-Event Bayesian Models. PLoS ONE, 2013, 8, e63240.	2.5	51
103	Ethnopharmacological evaluation of the informant consensus model on anti-tuberculosis claims among the Manus. Journal of Ethnopharmacology, 2006, 106, 82-89.	4.1	50
104	lleabethoxazole: a novel benzoxazole alkaloid with antimycobacterial activity. Tetrahedron Letters, 2006, 47, 3229-3232.	1.4	50
105	Indole alkaloids from two cultured cyanobacteria, Westiellopsis sp. and Fischerella muscicola. Bioorganic and Medicinal Chemistry, 2012, 20, 5290-5295.	3.0	50
106	Discovery and Characterization of the Tuberculosis Drug Lead Ecumicin. Organic Letters, 2014, 16, 6044-6047.	4.6	50
107	Strategies in anti-Mycobacterium tuberculosis drug discovery based on phenotypic screening. Journal of Antibiotics, 2019, 72, 719-728.	2.0	50
108	Sesquiterpenes from <i>Oplopanax horridus</i> . Journal of Natural Products, 2010, 73, 563-567.	3.0	49

#	Article	IF	Citations
109	Synthesis and evaluation of analogues of the tuberculosis drug bedaquiline containing heterocyclic B-ring units. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 5190-5196.	2.2	49
110	Oleanane Triterpenes from Junellia tridens. Journal of Natural Products, 2000, 63, 1611-1614.	3.0	48
111	Discovery and Development of the Covalent Hydrates of Trifluoromethylated Pyrazoles as Riboflavin Synthase Inhibitors with Antibiotic Activity Against <i>Mycobacterium tuberculosis</i> Organic Chemistry, 2009, 74, 5297-5303.	3.2	48
112	Identification, Synthesis, and Pharmacological Evaluation of Tetrahydroindazole Based Ligands as Novel Antituberculosis Agents. Journal of Medicinal Chemistry, 2010, 53, 649-659.	6.4	48
113	7-Substituted 2-Nitro-5,6-dihydroimidazo[2,1- $\langle i \rangle$ b $\langle j \rangle$ ][1,3]oxazines: Novel Antitubercular Agents Lead to a New Preclinical Candidate for Visceral Leishmaniasis. Journal of Medicinal Chemistry, 2017, 60, 4212-4233.	6.4	47
114	Inhibition of Mycobacterium tuberculosis Growth by Saringosterol from Lessoniani grescens. Journal of Natural Products, 2001, 64, 1463-1464.	3.0	46
115	Efficacy of Quinoxaline-2-Carboxylate 1,4-Di- N -Oxide Derivatives in Experimental Tuberculosis. Antimicrobial Agents and Chemotherapy, 2008, 52, 3321-3326.	3.2	46
116	Synthesis, Biological Evaluation, and Structureâ 'Activity Relationships for 5-[( <i>E</i> )-2-Arylethenyl]-3-isoxazolecarboxylic Acid Alkyl Ester Derivatives as Valuable Antitubercular Chemotypes. Journal of Medicinal Chemistry, 2009, 52, 6287-6296.	6.4	46
117	Antimycobacterial agents from selected Mexican medicinal plants. Journal of Pharmacy and Pharmacology, 2010, 57, 1117-1126.	2.4	46
118	Ruthenium (II) phosphine/picolinate complexes as antimycobacterial agents. European Journal of Medicinal Chemistry, 2010, 45, 598-601.	5 <b>.</b> 5	46
119	Repositioning Antitubercular 6-Nitro-2,3-dihydroimidazo[2,1-⟨i⟩b⟨/i⟩][1,3]oxazoles for Neglected Tropical Diseases: Structure–Activity Studies on a Preclinical Candidate for Visceral Leishmaniasis. Journal of Medicinal Chemistry, 2016, 59, 2530-2550.	6.4	46
120	Syntheses and Antituberculosis Activity of 1,3-Benzothiazinone Sulfoxide and Sulfone Derived from BTZ043. ACS Medicinal Chemistry Letters, 2015, 6, 128-133.	2.8	45
121	Novel sesquiterpenes and a lactone from the Jamaican sponge Myrmekioderma styx. Tetrahedron Letters, 2002, 43, 9699-9702.	1.4	44
122	Mycobacterium tuberculosis and cholinesterase inhibitors from Voacanga globosa. European Journal of Medicinal Chemistry, 2011, 46, 3118-3123.	5.5	43
123	Bioautography with TLC-MS/NMR for Rapid Discovery of Anti-tuberculosis Lead Compounds from Natural Sources. ACS Infectious Diseases, 2016, 2, 294-301.	3.8	43
124	Design, Synthesis, and Characterization of N-Oxide-Containing Heterocycles with in Vivo Sterilizing Antitubercular Activity. Journal of Medicinal Chemistry, 2017, 60, 8647-8660.	6.4	43
125	Inhibitory Effect of Sterols fromRuprechtia trifloraand Diterpenes fromCalceolaria pinnifoliaon the Growth ofMycobacterium tuberculosis. Planta Medica, 2003, 69, 628-631.	1.3	42
126	Synthesis and antitubercular activity of quaternized promazine and promethazine derivatives. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 1346-1348.	2.2	42

#	Article	IF	Citations
127	Carbamidocyclophanes F and G with anti-Mycobacterium tuberculosis activity from the cultured freshwater cyanobacterium Nostoc sp Tetrahedron Letters, 2014, 55, 686-689.	1.4	42
128	Development of $(6 < i > R < / i > )$ -2-Nitro-6-[4-(trifluoromethoxy)phenoxy]-6,7-dihydro-5 $< i > H < / i > -i midazo[2,1-< i > b < / i > ][1,3]oxazine (DNDI-8219): A New Lead for Visceral Leishmaniasis. Journal of Medicinal Chemistry, 2018, 61, 2329-2352.$	6.4	42
129	Antimycobacterial Activities of Dehydrocostus Lactone and Its Oxidation Products. Journal of Natural Products, 1998, 61, 1181-1186.	3.0	41
130	Antitubercular Activity and Inhibitory Effect on Epstein-Barr Virus Activation of Sterols and Polyisoprenepolyols from an Edible Mushroom, Hypsizigus marmoreus. Biological and Pharmaceutical Bulletin, 2005, 28, 1117-1119.	1.4	41
131	Structure–activity relationships of compounds targeting mycobacterium tuberculosis 1-deoxy-d-xylulose 5-phosphate synthase. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 5320-5323.	2.2	41
132	Succinylphosphonate Esters Are Competitive Inhibitors of MenD That Show Active-Site Discrimination between Homologous α-Ketoglutarate-Decarboxylating Enzymes. Biochemistry, 2010, 49, 2672-2679.	2.5	41
133	Synthesis and inÂvitro antimalarial and antitubercular activity of gold(III) complexes containing thiosemicarbazone ligands. Journal of Organometallic Chemistry, 2011, 696, 3392-3396.	1.8	41
134	Design, synthesis and anti-tubercular evaluation of new 2-acylated and 2-alkylated amino-5-(4-(benzyloxy)phenyl)thiophene-3-carboxylic acid derivatives. Part 1. European Journal of Medicinal Chemistry, 2011, 46, 3551-3563.	5.5	41
135	Mycobacterium tuberculosisGrowth Inhibition by Constituents ofSapiumhaematospermum. Journal of Natural Products, 2004, 67, 598-603.	3.0	40
136	Novel thiolactone–isatin hybrids as potential antimalarial and antitubercular agents. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 2055-2058.	2.2	40
137	Structure-activity relationships for unit C pyridyl analogues of the tuberculosis drug bedaquiline. Bioorganic and Medicinal Chemistry, 2019, 27, 1283-1291.	3.0	39
138	Development of an extraction method for mycobacterial metabolome analysis. Journal of Pharmaceutical and Biomedical Analysis, 2006, 41, 196-200.	2.8	38
139	Novel ring B abeo-sterols as growth inhibitors of Mycobacterium tuberculosis isolated from a Caribbean Sea sponge, Svenzea zeai. Tetrahedron Letters, 2007, 48, 8851-8854.	1.4	38
140	Antiâ€TB polyynes from the roots of <i>Angelica sinensis</i> . Phytotherapy Research, 2008, 22, 878-882.	5.8	38
141	Structural Basis for Catalysis of a Tetrameric Class IIa Fructose 1,6-Bisphosphate Aldolase from Mycobacterium tuberculosis. Journal of Molecular Biology, 2009, 386, 1038-1053.	4.2	38
142	Chlorinated Coumarins from the Polypore Mushroom <i>Fomitopsis officinalis</i> and Their Activity against <i>Mycobacterium tuberculosis</i> Journal of Natural Products, 2013, 76, 1916-1922.	3.0	38
143	Scaffold-switching: An exploration of 5,6-fused bicyclic heteroaromatics systems to afford antituberculosis activity akin to the imidazo[1,2-a]pyridine-3-carboxylates. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 3493-3498.	2.2	38
144	Putting Tuberculosis (TB) To Rest: Transformation of the Sleep Aid, Ambien, and "Anagrams―Generated Potent Antituberculosis Agents. ACS Infectious Diseases, 2015, 1, 85-90.	3.8	38

#	Article	IF	CITATIONS
145	Antimycobacterial Triterpenes from Melia volkensii. Journal of Natural Products, 1999, 62, 546-548.	3.0	37
146	Antitubercular Activity of Triterpenoids from Lippia turbinata. Journal of Natural Products, 2001, 64, 37-41.	3.0	37
147	Antitubercular triterpenes and phytosterols from Pandanus tectorius Soland. var. laevis. Journal of Natural Medicines, 2008, 62, 232-235.	2.3	37
148	Structure and Anti-TB Activity of Trachylobanes from the Liverwort <i>Jungermannia exsertifolia ssp. cordifolia </i> Journal of Natural Products, 2010, 73, 656-663.	3.0	37
149	Metal complexes of carboxamidrazone analogs as antitubercular agents. Journal of Inorganic Biochemistry, 2002, 90, 127-136.	3.5	36
150	Trypanoside, anti-tuberculosis, leishmanicidal, and cytotoxic activities of tetrahydrobenzothienopyrimidines. Bioorganic and Medicinal Chemistry, 2010, 18, 2880-2886.	3.0	36
151	Inhibition of <i>Mycobacterium tuberculosis</i> Methionine Aminopeptidases by Bengamide Derivatives. ChemMedChem, 2011, 6, 1041-1048.	3.2	36
152	Determinants of the Inhibition of DprE1 and CYP2C9 by Antitubercular Thiophenes. Angewandte Chemie - International Edition, 2017, 56, 13011-13015.	13.8	36
153	Antitubercular Nitroimidazoles Revisited: Synthesis and Activity of the Authentic 3-Nitro Isomer of Pretomanid. ACS Medicinal Chemistry Letters, 2017, 8, 1275-1280.	2.8	36
154	Current Prospects of Synthetic Curcumin Analogs and Chalcone Derivatives Against Mycobacterium Tuberculosis. Medicinal Chemistry, 2013, 9, 897-903.	1.5	35
155	Unusual antimalarial meroditerpenes from tropical red macroalgae. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 5662-5665.	2.2	34
156	Synthesis and characterization of pyruvate–isoniazid analogs and their copper complexes as potential ICL inhibitors. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 3172-3176.	2.2	34
157	Comparative in vitro antimicrobial activity of chinese medicinal herbs. Journal of Ethnopharmacology, 1986, 15, 279-288.	4.1	33
158	Design and Syntheses of Anti-Tuberculosis Agents Inspired by BTZ043 Using a Scaffold Simplification Strategy. ACS Medicinal Chemistry Letters, 2014, 5, 587-591.	2.8	33
159	Synthesis and Structure–Activity Relationships for Extended Side Chain Analogues of the Antitubercular Drug (6 <i>&gt;S</i> )-2-Nitro-6-{[4-(trifluoromethoxy)benzyl]oxy}-6,7-dihydro-5 <i>H</i> -imidazo[2,1- <i>b</i> ][1,3]oxazine (PA-824), Journal of Medicinal Chemistry, 2015, 58, 3036-3059.	6.4	33
160	New Phenylethanoids from Buddleja cordata subsp. cordata. Planta Medica, 2000, 66, 257-261.	1.3	32
161	Activity of Scottish Plant, Lichen and Fungal Endophyte Extracts against <i>Mycobacterium aurum</i> and <i>Mycobacterium tuberculosis</i> . Phytotherapy Research, 2010, 24, 692-698.	5.8	32
162	Synthesis of 3-(3-aryl-pyrrolidin-1-yl)-5-aryl-1,2,4-triazines that have antibacterial activity and also inhibit inorganic pyrophosphatase. Bioorganic and Medicinal Chemistry, 2014, 22, 406-418.	3.0	32

#	Article	IF	Citations
163	Diaza-anthracene Antibiotics from a Freshwater-Derived Actinomycete with Selective Antibacterial Activity toward <i>Mycobacterium tuberculosis</i> . ACS Infectious Diseases, 2015, 1, 168-174.	3.8	32
164	Antimycobacterial Matricaria Esters and Lactones from Astereae Species. Planta Medica, 1998, 64, 665-667.	1.3	31
165	Eucapsitrione, an Anti- <i>Mycobacterium tuberculosis</i> Anthraquinone Derivative from the Cultured Freshwater Cyanobacterium <i>Eucapsis</i> sp Journal of Natural Products, 2010, 73, 1441-1443.	3.0	31
166	Inhibiting enoyl-ACP reductase (Fabl) across pathogenic microorganisms by linear sesquiterpene lactones from Anthemis auriculata. Phytomedicine, 2008, 15, 1125-1129.	5.3	30
167	Phytoconstituents from Alpinia purpurata and their in vitro inhibitory activity against Mycobacterium tuberculosis. Pharmacognosy Magazine, 2010, 6, 339.	0.6	30
168	Antituberculosis Cycloartane Triterpenoids from <i>Radermachera boniana</i> . Journal of Natural Products, 2011, 74, 1318-1322.	3.0	30
169	Synthesis and antimycobacterial activities of non-purine analogs of 6-aryl-9-benzylpurines: Imidazopyridines, pyrrolopyridines, benzimidazoles, and indoles. Bioorganic and Medicinal Chemistry, 2011, 19, 3483-3491.	3.0	30
170	Potential of Lichen Secondary Metabolites against <i>Plasmodium</i> Liver Stage Parasites with FAS-II as the Potential Target. Journal of Natural Products, 2013, 76, 1064-1070.	3.0	30
171	In Vitro and In Vivo Activities of Ruthenium(II) Phosphine/Diimine/Picolinate Complexes (SCAR) against Mycobacterium tuberculosis. PLoS ONE, 2013, 8, e64242.	2.5	30
172	Syntheses and studies of quinolone-cephalosporins as potential anti-tuberculosis agents. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 5534-5537.	2.2	29
173	The design, synthesis, in silico ADME profiling, antiplasmodial and antimycobacterial evaluation of new arylamino quinoline derivatives. European Journal of Medicinal Chemistry, 2012, 57, 259-267.	5.5	29
174	(+) $\hat{a}$ Totarol from Chamaecyparis nootkatensis and activity against Mycobacterium tuberculosis. FA¬toterapìâ, 2001, 72, 572-574.	2.2	28
175	Antimycobacterial flavones from Haplopappus sonorensis. Fìtoterapìâ, 2003, 74, 226-230.	2.2	28
176	Advanced applications of counter-current chromatography in the isolation of anti-tuberculosis constituents from Dracaena angustifolia. Journal of Chromatography A, 2007, 1151, 169-174.	3.7	28
177	Fluorescence-based assay for polyprenyl phosphate-GlcNAc-1-phosphate transferase (WecA) and identification of novel antimycobacterial WecA inhibitors. Analytical Biochemistry, 2016, 512, 78-90.	2.4	28
178	Anti-tuberculosis activity and structure–activity relationships of oxygenated tricyclic carbazole alkaloids and synthetic derivatives. Bioorganic and Medicinal Chemistry, 2017, 25, 6167-6174.	3.0	28
179	Antimycobacterial Rufomycin Analogues from <i>Streptomyces atratus</i> Strain MJM3502. Journal of Natural Products, 2020, 83, 657-667.	3.0	28
180	Quantitative Purity–Activity Relationships of Natural Products: The Case of Anti-Tuberculosis Active Triterpenes from <i>Oplopanax horridus</i> . Journal of Natural Products, 2013, 76, 413-419.	3.0	27

#	Article	IF	CITATIONS
181	Diterpenes from Solidago rugosa. Phytochemistry, 1995, 38, 451-456.	2.9	26
182	Constituents of Seneciochionophilus with Potential Antitubercular Activity. Journal of Natural Products, 2004, 67, 1483-1487.	3.0	26
183	Antituberculotic and Antiprotozoal Activities of Primin, a Natural Benzoquinone:In vitro andin vivo Studies. Chemistry and Biodiversity, 2006, 3, 1230-1237.	2.1	26
184	Recent Advances in Methodologies for the Discovery of Antimycobacterial Drugs. Current Bioactive Compounds, 2007, 3, 201-208.	0.5	26
185	Synthesis and evaluation of rifabutin analogs against Mycobacterium avium and H37Rv, MDR and NRP Mycobacterium tuberculosis. Bioorganic and Medicinal Chemistry, 2009, 17, 503-511.	3.0	26
186	Tetrahydroxanthene-1,3( $2$ <i>H</i> )-dione Derivatives from <i>Uvaria valderramensis</i> . Journal of Natural Products, 2014, 77, 2711-2715.	3.0	26
187	6-Nitro-2,3-dihydroimidazo[2,1-b][1,3]thiazoles: Facile synthesis and comparative appraisal against tuberculosis and neglected tropical diseases. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 2583-2589.	2.2	26
188	Quinoline alkaloids from Lunasia amara inhibit Mycobacterium tuberculosis H37Rv in vitro. International Journal of Antimicrobial Agents, 2007, 29, 744-746.	2.5	25
189	Semisynthetic Studies on the Manzamine Alkaloids. Journal of Natural Products, 2008, 71, 300-308.	3.0	25
190	Syntheses of mycobactin analogs as potent and selective inhibitors of Mycobacterium tuberculosis. Organic and Biomolecular Chemistry, 2012, 10, 7584.	2.8	25
191	Antitubercular constituents from Premna odorata Blanco. Journal of Ethnopharmacology, 2014, 154, 471-474.	4.1	25
192	Imidazo[1,2- <i>a</i> ]Pyridine-3-Carboxamides Are Active Antimicrobial Agents against Mycobacterium avium Infection <i>In Vivo</i> Antimicrobial Agents and Chemotherapy, 2016, 60, 5018-5022.	3.2	25
193	Sweet spot matching: A thin-layer chromatography-based countercurrent solvent system selection strategy. Journal of Chromatography A, 2017, 1504, 46-54.	3.7	25
194	Variations in the C-unit of bedaquiline provides analogues with improved biology and pharmacology. Bioorganic and Medicinal Chemistry, 2020, 28, 115213.	3.0	25
195	2-Methoxylated fatty acids in marine sponges: Defense mechanism against mycobacteria?. Lipids, 2004, 39, 675-680.	1.7	24
196	HTS, Chemical Hybridization, and Drug Design Identify a Chemically Unique Antituberculosis Agent–Coupling Serendipity and Rational Approaches to Drug Discovery. ChemMedChem, 2007, 2, 811-813.	<b>3.</b> 2	24
197	Library Synthesis Using 5,6,7,8-Tetrahydro-1,6-naphthyridines as Scaffolds. ACS Combinatorial Science, 2008, 10, 534-540.	3.3	24
198	Residual Complexity Does Impact Organic Chemistry and Drug Discovery: The Case of Rufomyazine and Rufomycin. Journal of Organic Chemistry, 2018, 83, 6664-6672.	3.2	24

#	Article	IF	Citations
199	Evaluation of a modified antimycobacterial susceptibility test using Middlebrook 7H10 agar containing 2,3-diphenyl-5-thienyl-(2)-tetrazolium chloride. Journal of Microbiological Methods, 2006, 66, 548-551.	1.6	23
200	2,6-hexadecadiynoic acid and 2,6-nonadecadiynoic acid: Novel synthesized acetylenic fatty acids as potent antifungal agents. Lipids, 2006, 41, 507-511.	1.7	23
201	Natural product leads for drug discovery: Isolation, synthesis and biological evaluation of 6-cyano-5-methoxyindolo[2,3-a]carbazole based ligands as antibacterial agents. Bioorganic and Medicinal Chemistry, 2009, 17, 7126-7130.	3.0	23
202	Synthesis of non-purine analogs of 6-aryl-9-benzylpurines, and their antimycobacterial activities. Compounds modified in the imidazole ring. Bioorganic and Medicinal Chemistry, 2010, 18, 7274-7282.	3.0	23
203	Structure-Based Design of Novel Benzoxazinorifamycins with Potent Binding Affinity to Wild-Type and Rifampin-Resistant Mutant <i>Mycobacterium tuberculosis</i> RNA Polymerases. Journal of Medicinal Chemistry, 2012, 55, 3814-3826.	6.4	23
204	New finding of an anti-TB compound in the genus Marsypopetalum (Annonaceae) from a traditional herbal remedy of Laos. Journal of Ethnopharmacology, 2014, 151, 903-911.	4.1	23
205	Computer-aided discovery of two novel chalcone-like compounds active and selective against Leishmania infantum. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 2459-2464.	2.2	23
206	Identification of benzothiazinones containing an oxime functional moiety as new anti-tuberculosis agents. European Journal of Medicinal Chemistry, 2019, 181, 111595.	5.5	23
207	"Studies on Biodiversity of Vietnam and Laos―1998â^'2005: Examining the Impact#. Journal of Natural Products, 2006, 69, 473-481.	3.0	22
208	Dereplication of pentacyclic triterpenoids in plants by GC-EI/MS. Phytochemical Analysis, 2006, 17, 102-106.	2.4	22
209	Identification and Characterization of Novel Inhibitors of mPTPB, an Essential Virulent Phosphatase from Mycobacterium tuberculosis. ACS Medicinal Chemistry Letters, 2010, 1, 355-359.	2.8	22
210	Discovery and Optimization of Benzotriazine Di- <i>N</i> Oxides Targeting Replicating and Nonreplicating Mycobacterium tuberculosis. Journal of Medicinal Chemistry, 2012, 55, 6047-6060.	6.4	22
211	Construction and functionalization of fused pyridine ring leading to novel compounds as potential antitubercular agents. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 4629-4635.	2.2	22
212	In vitrosusceptibility of Mycobacterium tuberculosisto extracts of Eucalyptus camaldulensis and Eucalyptus to relliana and isolated compounds. Pharmaceutical Biology, 2012, 50, 92-98.	2.9	22
213	Trichormamides C and D, antiproliferative cyclic lipopeptides from the cultured freshwater cyanobacterium cf. Oscillatoria sp. UIC 10045. Bioorganic and Medicinal Chemistry, 2015, 23, 3153-3162.	3.0	22
214	New diterpenes of the pseudopterane class from two closely related Pseudopterogorgia species: isolation, structural elucidation, and biological evaluation. Tetrahedron, 2006, 62, 6998-7008.	1.9	21
215	Active Site Loop Dynamics of a Class Ila Fructose 1,6-Bisphosphate Aldolase from <i>Mycobacterium tuberculosis</i> . Biochemistry, 2013, 52, 912-925.	2.5	21
216	Synthesis and preliminary biological evaluation of a small library of hybrid compounds based on Ugi isocyanide multicomponent reactions with a marine natural product scaffold. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 5339-5343.	2.2	21

#	Article	IF	CITATIONS
217	Design, synthesis and evaluation of diarylpiperazine derivatives as potent anti-tubercular agents. European Journal of Medicinal Chemistry, 2015, 105, 238-244.	5.5	21
218	Phytochemical, Morphological, and Biological Investigations of Propolis from Central Chile. Zeitschrift Fur Naturforschung - Section C Journal of Biosciences, 1999, 54, 406-416.	1.4	20
219	An NMR method towards the routine chiral determination of natural products. Phytochemical Analysis, 2004, 15, 213-219.	2.4	20
220	Synthesis and in vitro biological evaluation of ring B abeo-sterols as novel inhibitors of Mycobacterium tuberculosis. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 5448-5450.	2.2	20
221	Derivatives of 3-Isoxazolecarboxylic Acid Esters - A Potent and Selective Compound Class against Replicating and Nonreplicating Mycobacterium tuberculosis. Current Topics in Medicinal Chemistry, 2012, 12, 729-734.	2.1	20
222	Natural product-based synthesis of novel anti-infective isothiocyanate- and isoselenocyanate-functionalized amphilectane diterpenes. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 854-857.	2.2	20
223	Antitubercular and Cytotoxic Chlorinated <i>seco</i> -Cyclohexenes from <i>Uvaria alba</i> . Journal of Natural Products, 2017, 80, 3319-3323.	3.0	19
224	Constituents of Quinchamalium majus with Potential Antitubercular Activity. Zeitschrift Fur Naturforschung - Section C Journal of Biosciences, 2004, 59, 797-802.	1.4	18
225	Structure-activity relationships of macrolides against Mycobacterium tuberculosis. Tuberculosis, 2008, 88, S49-S63.	1.9	18
226	glpX Gene of Mycobacterium tuberculosis: Heterologous Expression, Purification, and Enzymatic Characterization of the Encoded Fructose 1,6-bisphosphatase II. Applied Biochemistry and Biotechnology, 2011, 164, 1376-1389.	2.9	18
227	Hytramycins V and I, Anti-Mycobacterium tuberculosisHexapeptides from aStreptomyces hygroscopicusStrain. Journal of Natural Products, 2013, 76, 2009-2018.	3.0	18
228	Discovery of antitubercular 2,4-diphenyl-1H-imidazoles from chemical library repositioning and rational design. European Journal of Medicinal Chemistry, 2015, 100, 44-49.	5.5	18
229	Discovery of new leads against Mycobacterium tuberculosis using scaffold hopping and shape based similarity. Bioorganic and Medicinal Chemistry, 2017, 25, 4835-4844.	3.0	18
230	Quality Control of Therapeutic Peptides by <sup>1</sup> H NMR HiFSA Sequencing. Journal of Organic Chemistry, 2019, 84, 3055-3073.	3.2	18
231	Identification of Pyrazolo[1,5-a]pyridine-3-carboxamide Diaryl Derivatives as Drug Resistant Antituberculosis Agents. ACS Medicinal Chemistry Letters, 2019, 10, 295-299.	2.8	18
232	Design, synthesis and biological evaluation of novel 1,2,3-triazole analogues of Imidazo-[1,2-a]-pyridine-3-carboxamide against Mycobacterium tuberculosis. Toxicology in Vitro, 2021, 74, 105137.	2.4	18
233	A new variant of Autographa californica nuclear polyhedrosis virus. Journal of Invertebrate Pathology, 1980, 36, 159-165.	3.2	17
234	NMR and Molecular Mechanics Study of Pyrethrins I and II. Journal of Agricultural and Food Chemistry, 1999, 47, 3402-3410.	5.2	17

#	Article	IF	CITATIONS
235	New C-3 $\hat{a}$ e $^2$ hydroxamate-substituted and more lipophilic cyclic hydroxamate cephalosporin derivatives as a potential new generation of selective antimicrobial agents. Organic and Biomolecular Chemistry, 2006, 4, 4178-4185.	2.8	17
236	Biological evaluation of plants of Laos used in the treatment of tuberculosis in Lao traditional medicine. Pharmaceutical Biology, 2009, 47, 26-33.	2.9	17
237	Allylic thiocyanates as a new class of antitubercular agents. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 6486-6489.	2.2	17
238	Heteroaryl ether analogues of an antileishmanial 7-substituted 2-nitroimidazooxazine lead afford attenuated hERG risk: InÂvitro and inÂvivo appraisal. European Journal of Medicinal Chemistry, 2021, 209, 112914.	5.5	17
239	Pyrazole and imidazo[1,2-b]pyrazole Derivatives as New Potential Antituberculosis Agents. Medicinal Chemistry, 2019, 15, 17-27.	1.5	17
240	Identification of heteroarylenamines as a new class of antituberculosis lead molecules. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 4097-4099.	2.2	16
241	Antitubercular sterols fromThalia multiflora Horkel ex Koernicke. Phytotherapy Research, 2005, 19, 876-880.	5.8	16
242	Utilization of the Suzuki Coupling to Enhance the Antituberculosis Activity of Aryloxazoles. Heterocycles, 2010, 80, 977.	0.7	16
243	Airborne Antituberculosis Activity of <i>Eucalyptus citriodora</i> Products, 2014, 77, 603-610.	3.0	16
244	Synthesis and structure-activity relationships for tetrahydroisoquinoline-based inhibitors of Mycobacterium tuberculosis. Bioorganic and Medicinal Chemistry, 2020, 28, 115784.	3.0	16
245	Antitubercular activity of the semi–polar extractives of Uvaria rufa. Asian Pacific Journal of Tropical Medicine, 2012, 5, 777-780.	0.8	15
246	Design, synthesis, and evaluation of 4-(substituted)phenyl-2-thioxo-3,4-dihydro-1H-chromino[4,3-d]pyrimidin-5-one and 4-(substituted)phenyl-3,4-dihydro-1H-chromino[4,3-d]pyrimidine-2,5-dione analogs as antitubercular agents. Medicinal Chemistry Research, 2014, 23, 2564-2575.	2.4	15
247	Cytotoxic Constituents from <i>Lobaria scrobiculata</i> and a Comparison of Two Bioassays for Their Evaluation. Journal of Natural Products, 2014, 77, 1069-1073.	3.0	15
248	Mycobacterial Plasmids. Microbiology and Immunology, 1986, 30, 903-907.	1.4	14
249	Axenic incorporation of [U-14C]palmitic acid into the phenolic glycolipid-I ofMycobacterium leprae. FEMS Microbiology Letters, 1987, 48, 407-411.	1.8	14
250	The Latin American ICBG: The First Five Years. Pharmaceutical Biology, 1999, 37, 35-54.	2.9	14
251	Modification of the side chain of micromolide, an anti-tuberculosis natural product. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 5311-5315.	2.2	13
252	6â€Hydrogenâ€8â€Methylquinolones Active Against Replicating and Nonâ€replicating <i>Mycobacterium tuberculosis</i> . Chemical Biology and Drug Design, 2012, 80, 781-786.	3.2	13

#	Article	IF	CITATIONS
253	Isolation of Tryptanthrin and Reassessment of Evidence for Its Isobaric Isostere Wrightiadione in Plants of the Wrightia Genus. Journal of Natural Products, 2019, 82, 440-448.	3.0	13
254	Discovery and preclinical profile of sudapyridine (WX-081), a novel anti-tuberculosis agent. Bioorganic and Medicinal Chemistry Letters, 2022, 71, 128824.	2.2	13
255	CCC in the Phytochemical Analysis of Antiâ€Tuberculosis Ethnobotanicals. Journal of Liquid Chromatography and Related Technologies, 2005, 28, 2017-2028.	1.0	12
256	Release of nitrite from the antitubercular nitroimidazoledrug PA-824 and analogues upon one-electron reduction in protic, non-aqueous solvent. Organic and Biomolecular Chemistry, 2010, 8, 413-418.	2.8	12
257	Biarylmethoxy 2-nitroimidazooxazine antituberculosis agents: Effects of proximal ring substitution and linker reversal on metabolism and efficacy. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 3804-3809.	2.2	12
258	Design, syntheses, and anti-tuberculosis activities of conjugates of piperazino-1,3-benzothiazin-4-ones (pBTZs) with 2,7-dimethylimidazo [1,2-a]pyridine-3-carboxylic acids and 7-phenylacetyl cephalosporins. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 2068-2071.	2.2	12
259	Syntheses and biological evaluations of highly functionalized hydroxamate containing and $\langle i \rangle N \langle i \rangle$ -methylthio monobactams as anti-tuberculosis and $\hat{l}^2$ -lactamase inhibitory agents. MedChemComm, 2016, 7, 141-147.	3.4	12
260	Synthesis and Activity against Mycobacterium tuberculosis of Olivacine and Oxygenated Derivatives. Molecules, 2018, 23, 1402.	3.8	12
261	An antimycobacterial pleuromutilin analogue effective against dormant bacilli. Bioorganic and Medicinal Chemistry, 2018, 26, 4787-4796.	3.0	12
262	Mce3R Stress-Resistance Pathway Is Vulnerable to Small-Molecule Targeting That Improves Tuberculosis Drug Activities. ACS Infectious Diseases, 2019, 5, 1239-1251.	3.8	12
263	Antitubercular polyhalogenated phenothiazines and phenoselenazine with reduced binding to CNS receptors. European Journal of Medicinal Chemistry, 2020, 201, 112420.	5.5	12
264	glpx Gene in Mycobacterium tuberculosis Is Required for In Vitro Gluconeogenic Growth and In Vivo Survival. PLoS ONE, 2015, 10, e0138436.	2.5	12
265	NOC Chemistry for Tuberculosis—Further Investigations on the Structure–Activity Relationships of Antitubercular Isoxazoleâ€3â€carboxylic Acid Ester Derivatives. ChemMedChem, 2010, 5, 1667-1672.	3.2	11
266	Synthesis and structure–activity relationships of novel substituted 8-amino, 8-thio, and 1,8-pyrazole congeners of antitubercular rifamycin S and rifampin. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 6094-6099.	2.2	11
267	Syntheses and evaluation of substituted aromatic hydroxamates and hydroxamic acids that target Mycobacterium tuberculosis. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 4933-4936.	2.2	11
268	Anti-Mycobacterium tuberculosis Activity of Esters of Quinoxaline 1,4-Di-N-Oxide. Molecules, 2018, 23, 1453.	3.8	11
269	Induction of pyruvate decarboxylase in Candida utilis. Mycopathologia, 1983, 83, 29-33.	3.1	10
270	Chemical constituents from Xylosma longifolia and their anti-tubercular activity. Phytochemistry Letters, 2011, 4, 250-253.	1.2	10

#	Article	IF	CITATIONS
271	Bioassayâ€Guided Isolation and Structural Modification of the Antiâ€ <scp>TB</scp> Resorcinols from <i>Ardisia gigantifolia</i> . Chemical Biology and Drug Design, 2016, 88, 293-301.	3.2	10
272	An iboga alkaloid chemotaxonomic marker from endemic <i>Tabernaemontana ternifolia</i> with antitubercular activity. Natural Product Research, 2020, 34, 1175-1179.	1.8	10
273	Rufomycins or Ilamycins: Naming Clarifications and Definitive Structural Assignments. Journal of Natural Products, 2021, 84, 2644-2663.	3.0	10
274	Synthetic studies towards isomeric pyrazolopyrimidines as potential ATP synthesis inhibitors of Mycobacterium tuberculosis. Structural correction of reported N-(6-(2-(dimethylamino)ethoxy)-5-fluoropyridin-3-yl)-2-(4-fluorophenyl)-5-(trifluoromethyl)pyrazolo[1,5-α]pyrimidi Tetrahedron Letters, 2022, 90, 153611.	in-7-amine	.10
275	A novel indigoid anti-tuberculosis agent. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 268-270.	2.2	9
276	Structural Sequencing of Oligopeptides Aided by <sup>1</sup> H Iterative Full-Spin Analysis. Journal of Natural Products, 2017, 80, 2630-2643.	3.0	9
277	Attenuation of Mycobacterium species through direct and macrophage mediated pathway by unsymmetrical diaryl urea. European Journal of Medicinal Chemistry, 2017, 125, 825-841.	5.5	9
278	Antitubercular and cytotoxic polyoxygenated cyclohexane derivatives from <i>Uvaria grandiflora</i> . Natural Product Research, 2021, 35, 5229-5232.	1.8	9
279	Novel Linker Variants of Antileishmanial/Antitubercular 7-Substituted 2-Nitroimidazooxazines Offer Enhanced Solubility. ACS Medicinal Chemistry Letters, 2021, 12, 275-281.	2.8	9
280	<i>In Vitro</i> Profiling of Antitubercular Compounds by Rapid, Efficient, and Nondestructive Assays Using Autoluminescent Mycobacterium tuberculosis. Antimicrobial Agents and Chemotherapy, 2021, 65, e0028221.	3.2	9
281	A lanostane aldehyde from Momordica charantia. Phytochemistry Letters, 2012, 5, 682-684.	1.2	8
282	Syntheses and biological studies of novel spiropiperazinyl oxazolidinone antibacterial agents using a spirocyclic diene derived acylnitroso Dielsâ 'Alder reaction. Bioorganic and Medicinal Chemistry, 2012, 20, 3422-3428.	3.0	8
283	Rapid determination of growth inhibition of Mycobacterium tuberculosis by GC–MS/MS quantitation of tuberculostearic acid. Tuberculosis, 2013, 93, 322-329.	1.9	8
284	Benzylsulfanyl benzo-heterocycle amides and hydrazones as new agents against drug-susceptible and resistant Mycobacterium tuberculosis. MedChemComm, 2017, 8, 1303-1306.	3.4	8
285	Synthesis, antibacterial, and antitubercular studies of some novel isatin derivatives. Medicinal Chemistry Research, 2012, 21, 4335-4340.	2.4	7
286	Structural requirements for the antitubercular quaternized triflupromazine pharmacophore. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 5679-5680.	2.2	7
287	Antimycobacterial activity of pyrazinoate prodrugs in replicating and non-replicating Mycobacterium tuberculosis. Tuberculosis, 2016, 99, 11-16.	1.9	7
288	Exploring the Sponge Consortium <i>Plakortis symbiotica–Xestospongia deweerdtae</i> as a Potential Source of Antimicrobial Compounds and Probing the Pharmacophore for Antituberculosis Activity of Smenothiazole A by Diverted Total Synthesis. Journal of Natural Products, 2017, 80, 2295-2303.	3.0	7

#	Article	IF	Citations
289	Use of green fluorescent protein labeled non-tuberculous mycobacteria to evaluate the activity quaternary ammonium compound disinfectants and antibiotics. Brazilian Journal of Microbiology, 2017, 48, 151-158.	2.0	7
290	In Vitro Activities of Enantiopure and Racemic $1\hat{a}\in^2$ -Acetoxychavicol Acetate against Clinical Isolates of Mycobacterium tuberculosis. Scientia Pharmaceutica, 2017, 85, 32.	2.0	7
291	Synthesis and antimicrobial activities of N6-hydroxyagelasine analogs and revision of the structure of ageloximes. Bioorganic and Medicinal Chemistry, 2019, 27, 620-629.	3.0	7
292	Synthesis and structure-activity relationships for a new class of tetrahydronaphthalene amide inhibitors of Mycobacterium tuberculosis. European Journal of Medicinal Chemistry, 2022, 229, 114059.	5.5	7
293	Terpenes from Liatris ohlingerae. Phytochemistry, 1994, 37, 1295-1299.	2.9	6
294	Dihydroparthenolide diol, a novel sesquiterpene lactone. Acta Crystallographica Section E: Structure Reports Online, 2001, 57, o323-o325.	0.2	6
295	Vaccination with Bacilleâ€Calmette Guérin Promotes Mycobacterial Control in Guinea Pig Macrophages Infected In Vivo. Journal of Infectious Diseases, 2008, 198, 768-771.	4.0	6
296	Anti-tuberculosis Compounds from two Bolivian Medicinal Plants, Senecio Mathewsii and Usnea Florida. Natural Product Communications, 2008, 3, 1934578X0800300.	0.5	6
297	Crystallization and preliminary X-ray characterization of theglpX-encoded class II fructose-1,6-bisphosphatase fromMycobacterium tuberculosis. Acta Crystallographica Section F: Structural Biology Communications, 2011, 67, 710-713.	0.7	6
298	A novel combinatorial biocatalytic approach for producing antibacterial compounds effective against Mycobacterium tuberculosis (TB). Applied Microbiology and Biotechnology, 2013, 97, 7151-7163.	3.6	6
299	Synthesis and Evaluation as Antitubercular Agents of 5â€Arylethenyl and 5â€(Hetero)arylâ€3â€Isoxazolecarboxylate. Drug Development Research, 2013, 74, 162-172.	2.9	6
300	Biological Profiling Enables Rapid Mechanistic Classification of Phenotypic Screening Hits and Identification of KatG Activation-Dependent Pyridine Carboxamide Prodrugs With Activity Against Mycobacterium tuberculosis. Frontiers in Cellular and Infection Microbiology, 2020, 10, 582416.	3.9	6
301	2-Aryl benzazole derived new class of anti-tubercular compounds: Endowed to eradicate mycobacterium tuberculosis in replicating and non-replicating forms. Bioorganic Chemistry, 2020, 103, 104170.	4.1	5
302	1,3-Oxazine-2-one derived dual-targeted molecules against replicating and non-replicating forms of Mycobacterium tuberculosis. European Journal of Medicinal Chemistry, 2020, 208, 112835.	5.5	5
303	Insights into the Chemical Diversity of Selected Fungi from the Tza Itz $\tilde{A}_i$ Cenote of the Yucatan Peninsula. ACS Omega, 2022, 7, 12171-12185.	3.5	5
304	Induction of fermentation in Crabtree-negative yeasts. Mycopathologia, 1983, 82, 185-190.	3.1	4
305	Inhibitory effect of oxygenated cholestan- $3\hat{l}^2$ -ol derivatives on the growth of Mycobacterium tuberculosis. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 6111-6113.	2.2	4
306	Biophysical Screening of a Focused Library for the Discovery of CYP121 Inhibitors as Novel Antimycobacterials. ChemMedChem, 2017, 12, 1616-1626.	3.2	4

#	Article	IF	CITATIONS
307	Hydride-induced Meisenheimer complex formation reflects activity of nitro aromatic anti-tuberculosis compounds. RSC Medicinal Chemistry, 2021, 12, 62-72.	3.9	4
308	Design of Novel Phosphopantetheine Adenylyltransferase Inhibitors: A Potential New Approach to Tackle Mycobacterium tuberculosis. Current Topics in Medicinal Chemistry, 2021, 21, 1186-1197.	2.1	4
309	Chemical Diversity and Antimicrobial Potential of Cultivable Fungi from Deep-Sea Sediments of the Gulf of Mexico. Molecules, 2021, 26, 7328.	3.8	4
310	Villarinol, a new Alkenoyloxyalkenol Derivative from the Endemic Philippine Rubiaceae species Villaria odorata. Natural Product Communications, 2012, 7, 1934578X1200700.	0.5	3
311	Design, Synthesis and Antitubercular Evaluation of New 2-amino-5-(4-) Tj ETQq1 1 0.784314 rgBT /Overlock 10 Tf Discovery, 2014, 12, 29-37.	50 587 To 0.7	d ((benzylox 3
312	Rufomycin Exhibits Dual Effects Against Mycobacterium abscessus Infection by Inducing Host Defense and Antimicrobial Activities. Frontiers in Microbiology, 2021, 12, 695024.	3.5	3
313	Villarinol, a new alkenoyloxyalkenol derivative from the endemic Philippine Rubiaceae species Villaria odorata. Natural Product Communications, 2012, 7, 779-80.	0.5	3
314	Discovery and preclinical evaluations of JBD0131, a novel nitrodihydro-imidazooxazole anti-tuberculosis agent. Bioorganic and Medicinal Chemistry Letters, 2022, 72, 128871.	2.2	3
315	Quinolineâ€Proline, Triazole Hybrids: Design, Synthesis, Antituberculosis, Molecular Docking, and ADMET Studies. Journal of Heterocyclic Chemistry, 2021, 58, 952-968.	2.6	2
316	Determinants of the Inhibition of DprE1 and CYP2C9 by Antitubercular Thiophenes. Angewandte Chemie, 2017, 129, 13191-13195.	2.0	1
317	One-Pot Synthesis of Novel Hydrazono-1,3-Thıazolıdın-4-One Derivatives as Anti-HIV and Anti-Tubercular Agents: Synthesıs, Bıologıcal Evaluatıon, Molecular Modelling and Admet Studıes. Current HIV Research, 2022, 20, 255-271.	0.5	1
318	Optimization of Benzoxazinorifamycins to Minimize hPXR Activation for the Treatment of Tuberculosis and HIV Coinfection. ACS Infectious Diseases, 2022, 8, 1408-1421.	3.8	1
319	Optimization of Benzoxazinorifamycins to Improve <i>Mycobacterium tuberculosis</i> RNA Polymerase Inhibition and Treatment of Tuberculosis. ACS Infectious Diseases, 2022, 8, 1422-1438.	3.8	1
320	Clinical Trial of Sparfloxacin in the Treatment of Leprosy. Drugs, 1993, 45, 225-226.	10.9	0
321	Carcinogenic effects of N-nitroso-3-(substituted phenylimino)-indolin-2-one derivatives. Journal of Pharmacy and Bioallied Sciences, 2012, 4, 207.	0.6	O
322	Erratum to "A lanostane aldehyde from Momordica charantia―[Phytochem. Lett. 5 (2012) 682–684]. Phytochemistry Letters, 2012, 5, 819.	1.2	0
323	Photoactivated [3+2] Addition of 6,7- <i>seco</i> -angustilobine B to Fullerene [C <sub>60</sub> ]. Natural Product Communications, 2012, 7, 1934578X1200700.	0.5	O
324	A Potentially New Treatment for Tuberculosis; Will a Diarylquinoline Work for Leprosy?. International Journal of Leprosy and Other Mycobacterial Diseases, 2005, 73, 32.	0.3	0

#	ŧ	Article	lF	CITATIONS
3	25	Antiâ€tuberculosis Drug Discovery from Phenotypic Highâ€throughput Screening of Actinomycete Cultures. FASEB Journal, 2018, 32, lb633.	0.5	O
3	26	New Terpenoids from the Corticioid Fungus Punctularia atropurpurascens and their Antimycobacterial Evaluation. Planta Medica, 2022, , .	1.3	O