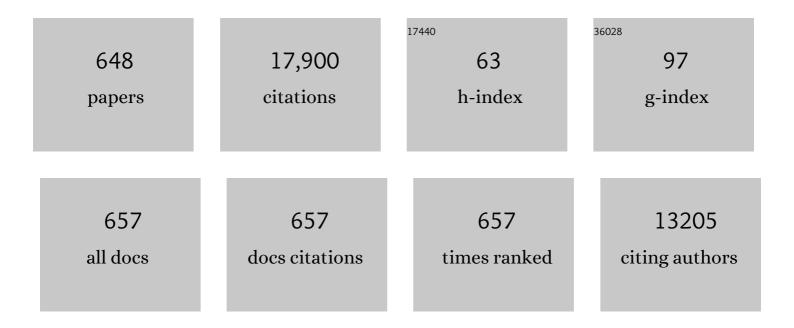
List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Acanthamoeba: biology and increasing importance in human health. FEMS Microbiology Reviews, 2006, 30, 564-595.	8.6	660
2	An update on <i>Acanthamoeba</i> keratitis: diagnosis, pathogenesis and treatment. Parasite, 2015, 22, 10.	2.0	494
3	Biology and pathogenesis of Acanthamoeba. Parasites and Vectors, 2012, 5, 6.	2.5	416
4	Load forecasting, dynamic pricing and DSM in smart grid: A review. Renewable and Sustainable Energy Reviews, 2016, 54, 1311-1322.	16.4	322
5	Schiff bases in medicinal chemistry: a patent review (2010-2015). Expert Opinion on Therapeutic Patents, 2017, 27, 63-79.	5.0	208
6	Biscoumarin: new class of urease inhibitors; economical synthesis and activity. Bioorganic and Medicinal Chemistry, 2004, 12, 1963-1968.	3.0	201
7	Tracking Five Millennia of Horse Management with Extensive Ancient Genome Time Series. Cell, 2019, 177, 1419-1435.e31.	28.9	195
8	Quinazoline and quinazolinone as important medicinal scaffolds: a comparative patent review (2011–2016). Expert Opinion on Therapeutic Patents, 2018, 28, 281-297.	5.0	165
9	Pathogenesis of Acanthamoeba infections. Microbial Pathogenesis, 2003, 34, 277-285.	2.9	162
10	Synthesis of novel inhibitors of α-glucosidase based on the benzothiazole skeleton containing benzohydrazide moiety and their molecular docking studies. European Journal of Medicinal Chemistry, 2015, 92, 387-400.	5.5	155
11	Pathogenesis of microbial keratitis. Microbial Pathogenesis, 2017, 104, 97-109.	2.9	155
12	Isatin based Schiff bases as inhibitors of α-glucosidase: Synthesis, characterization, in vitro evaluation and molecular docking studies. Bioorganic Chemistry, 2015, 60, 42-48.	4.1	147
13	Cytotoxic Necrotizing Factor-1 Contributes to Escherichia coli K1 Invasion of the Central Nervous System. Journal of Biological Chemistry, 2002, 277, 15607-15612.	3.4	145
14	Acanthamoeba genotype T4 from the UK and Iran and isolation of the T2 genotype from clinical isolates. Journal of Medical Microbiology, 2005, 54, 755-759.	1.8	139
15	Synthesis of bis-Schiff bases of isatins and their antiglycation activity. Bioorganic and Medicinal Chemistry, 2009, 17, 7795-7801.	3.0	134
16	Combined emission economic dispatch of power system including solar photo voltaic generation. Energy Conversion and Management, 2015, 92, 82-91.	9.2	129
17	Synthesis and molecular docking studies of potent α-glucosidase inhibitors based on biscoumarin skeleton. European Journal of Medicinal Chemistry, 2014, 81, 245-252.	5.5	128
18	Biology and pathogenesis of Naegleria fowleri. Acta Tropica, 2016, 164, 375-394.	2.0	127

#	Article	IF	CITATIONS
19	Triazinoindole analogs as potent inhibitors of α-glucosidase: Synthesis, biological evaluation and molecular docking studies. Bioorganic Chemistry, 2015, 58, 81-87.	4.1	126
20	Increasing Importance of <i>Balamuthia mandrillaris</i> . Clinical Microbiology Reviews, 2008, 21, 435-448.	13.6	121
21	Multicomponent reactions (MCR) in medicinal chemistry: a patent review (2010-2020). Expert Opinion on Therapeutic Patents, 2021, 31, 267-289.	5.0	115
22	Proteases as Markers for Differentiation of Pathogenic and Nonpathogenic Species of <i>Acanthamoeba</i> . Journal of Clinical Microbiology, 2000, 38, 2858-2861.	3.9	113
23	Escherichia coli K1 RS218 Interacts with Human Brain Microvascular Endothelial Cells via Type 1 Fimbria Bacteria in the Fimbriated State. Infection and Immunity, 2005, 73, 2923-2931.	2.2	112
24	Synthesis, in vitro evaluation and molecular docking studies of thiazole derivatives as new inhibitors of α-glucosidase. Bioorganic Chemistry, 2015, 62, 15-21.	4.1	109
25	Tetraketones: A new class of tyrosinase inhibitors. Bioorganic and Medicinal Chemistry, 2006, 14, 344-351.	3.0	99
26	Acanthamoeba interactions with human brain microvascular endothelial cells. Microbial Pathogenesis, 2003, 35, 235-241.	2.9	98
27	Synthesis and inÂvitro urease inhibitory activity of N,N′-disubstituted thioureas. European Journal of Medicinal Chemistry, 2014, 74, 314-323.	5.5	98
28	Antimicrobial activities of green synthesized gums-stabilized nanoparticles loaded with flavonoids. Scientific Reports, 2019, 9, 3122.	3.3	96
29	Pathogenicity, Morphology, and Differentiation of Acanthamoeba. Current Microbiology, 2001, 43, 391-395.	2.2	95
30	Primary Amoebic Meningoencephalitis Caused by Naegleria fowleri: An Old Enemy Presenting New Challenges. PLoS Neglected Tropical Diseases, 2014, 8, e3017.	3.0	95
31	Acanthamoeba castellanii Induces Host Cell Death via a Phosphatidylinositol 3-Kinase-Dependent Mechanism. Infection and Immunity, 2005, 73, 2704-2708.	2.2	94
32	Oxazolones: New tyrosinase inhibitors; synthesis and their structure–activity relationships. Bioorganic and Medicinal Chemistry, 2006, 14, 6027-6033.	3.0	93
33	Schiff bases of 3-formylchromone as thymidine phosphorylase inhibitors. Bioorganic and Medicinal Chemistry, 2009, 17, 2983-2988.	3.0	93
34	Escherichia coli interactions with Acanthamoeba: a symbiosis with environmental and clinical implications. Journal of Medical Microbiology, 2006, 55, 689-694.	1.8	91
35	Synthesis of novel inhibitors of β-glucuronidase based on benzothiazole skeleton and study of their binding affinity by molecular docking. Bioorganic and Medicinal Chemistry, 2011, 19, 4286-4294.	3.0	91
36	Determination of free phenolic acids and antioxidant activity of methanolic extracts obtained from fruits and leaves of Chenopodium album. Food Chemistry, 2011, 126, 1850-1855.	8.2	89

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37	Synthesis, α-glucosidase inhibition and molecular docking study of coumarin based derivatives. Bioorganic Chemistry, 2018, 77, 586-592.	4.1	88
38	Molecular and Physiological Differentiation Between Pathogenic and Nonpathogenic Acanthamoeba. Current Microbiology, 2002, 45, 197-202.	2.2	87
39	Acanthamoeba Can Be Differentiated by the Polymerase Chain Reaction and Simple Plating Assays. Current Microbiology, 2001, 43, 204-208.	2.2	86
40	Brain-Eating Amoebae: Silver Nanoparticle Conjugation Enhanced Efficacy of Anti-Amoebic Drugs against <i>Naegleria fowleri</i> . ACS Chemical Neuroscience, 2017, 8, 2626-2630.	3.5	85
41	Synthesis and in vitro acetylcholinesterase and butyrylcholinesterase inhibitory potential of hydrazide based Schiff bases. Bioorganic Chemistry, 2016, 68, 30-40.	4.1	82
42	Synthesis of novel derivatives of oxindole, their urease inhibition and molecular docking studies. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 3285-3289.	2.2	79
43	The Development of Drugs against Acanthamoeba Infections. Antimicrobial Agents and Chemotherapy, 2016, 60, 6441-6450.	3.2	79
44	FimH-mediated Escherichia coli K1 invasion of human brain microvascular endothelial cells. Cellular Microbiology, 2007, 9, 169-178.	2.1	78
45	Synthesis of diethyl 4-substituted-2,6-dimethyl-1,4-dihydropyridine-3,5-dicarboxylates as a new series of inhibitors against yeast α-glucosidase. European Journal of Medicinal Chemistry, 2015, 95, 199-209.	5.5	78
46	Syntheses of new 3-thiazolyl coumarin derivatives, inÂvitro α -glucosidase inhibitory activity, and molecular modeling studies. European Journal of Medicinal Chemistry, 2016, 122, 196-204.	5.5	78
47	Synthesis of benzotriazoles derivatives and their dual potential as α-amylase and α-glucosidase inhibitors inÂvitro: Structure-activity relationship, molecular docking, and kinetic studies. European Journal of Medicinal Chemistry, 2019, 183, 111677.	5.5	78
48	Outer membrane protein A and cytotoxic necrotizing factor-1 use diverse signaling mechanisms for Escherichia coli K1 invasion of human brain microvascular endothelial cells. Microbial Pathogenesis, 2003, 35, 35-42.	2.9	77
49	In Vitro Pathogenicity ofAcanthamoebals Associated with the Expression of the Mannose-Binding Protein. , 2006, 47, 1056.		76
50	Brain-Eating Amoebae: Predilection Sites in the Brain and Disease Outcome. Journal of Clinical Microbiology, 2017, 55, 1989-1997.	3.9	76
51	Synthesis of Coumarin Derivatives with Cytotoxic, Antibacterial and Antifungal Activity. Journal of Enzyme Inhibition and Medicinal Chemistry, 2004, 19, 373-379.	5.2	75
52	5-Bromo-2-aryl benzimidazole derivatives as non-cytotoxic potential dual inhibitors of α -glucosidase and urease enzymes. Bioorganic Chemistry, 2017, 72, 21-31.	4.1	75
53	Biology-oriented drug synthesis (BIODS) of 2-(2-methyl-5-nitro-1H-imidazol-1-yl)ethyl aryl ether derivatives, in vitro α-amylase inhibitory activity and in silico studies. Bioorganic Chemistry, 2017, 74, 1-9.	4.1	75
54	Acanthamoeba affects the integrity of human brain microvascular endothelial cells and degrades the tight junction proteins. International Journal for Parasitology, 2009, 39, 1611-1616.	3.1	73

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55	Carbohydrate analysis of Acanthamoeba castellanii. Experimental Parasitology, 2009, 122, 338-343.	1.2	72
56	Synthesis, α -glucosidase inhibitory activity and in silico study of tris -indole hybrid scaffold with oxadiazole ring: As potential leads for the management of type-II diabetes mellitus. Bioorganic Chemistry, 2017, 74, 30-40.	4.1	72
57	Synthesis, molecular docking and α-glucosidase inhibition of 5-aryl-2-(6′-nitrobenzofuran-2′-yl)-1,3,4-oxadiazoles. Bioorganic Chemistry, 2016, 66, 117-123.	4.1	71
58	Synthesis of Novel Bisindolylmethane Schiff bases and Their Antibacterial Activity. Molecules, 2014, 19, 11722-11740.	3.8	70
59	New Hybrid Hydrazinyl Thiazole Substituted Chromones: As Potential α-Amylase Inhibitors and Radical (DPPH & ABTS) Scavengers. Scientific Reports, 2017, 7, 16980.	3.3	70
60	High entropy alloy thin films of AlCoCrCu0.5FeNi with controlled microstructure. Applied Surface Science, 2019, 495, 143560.	6.1	69
61	Leptospirosis: Increasing importance in developing countries. Acta Tropica, 2020, 201, 105183.	2.0	68
62	Synthesis of new oxadiazole derivatives as α-glucosidase inhibitors. Bioorganic and Medicinal Chemistry, 2015, 23, 4155-4162.	3.0	67
63	Synthesis, β-glucuronidase inhibition and molecular docking studies of hybrid bisindole-thiosemicarbazides analogs. Bioorganic Chemistry, 2016, 68, 56-63.	4.1	66
64	Balamuthia amoebic encephalitis: An emerging disease with fatal consequences. Microbial Pathogenesis, 2008, 44, 89-97.	2.9	65
65	Synthesis, biological evaluation and molecular docking of N-phenyl thiosemicarbazones as urease inhibitors. Bioorganic Chemistry, 2015, 61, 51-57.	4.1	65
66	Hydrazinyl arylthiazole based pyridine scaffolds: Synthesis, structural characterization, inÂvitro α-glucosidase inhibitory activity, and in silico studies. European Journal of Medicinal Chemistry, 2017, 138, 255-272.	5.5	65
67	Synthesis, in vitro α-glucosidase inhibitory potential and molecular docking study of thiadiazole analogs. Bioorganic Chemistry, 2018, 78, 201-209.	4.1	65
68	Post-mortem culture of Balamuthia mandrillaris from the brain and cerebrospinal fluid of a case of granulomatous amoebic meningoencephalitis, using human brain microvascular endothelial cells. Journal of Medical Microbiology, 2004, 53, 1007-1012.	1.8	64
69	Synthesis, <i>In vitro</i> and Docking Studies of New Flavone Ethers as <i>α</i> â€Glucosidase Inhibitors. Chemical Biology and Drug Design, 2016, 87, 361-373.	3.2	63
70	Synthesis and β-glucuronidase inhibitory activity of 2-arylquinazolin-4(3H)-ones. Bioorganic and Medicinal Chemistry, 2014, 22, 3449-3454.	3.0	61
71	The Evolutionary Origin and Genetic Makeup of Domestic Horses. Genetics, 2016, 204, 423-434.	2.9	61
72	Extracellular proteases of (encephalitis isolate belonging to T1 genotype) contribute to increased permeability in an in vitro model of the human bloodâ€″brain barrier. Journal of Infection, 2005, 51, 150-156.	3.3	60

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73	Cellulose degradation: a therapeutic strategy in the improved treatment of Acanthamoeba infections. Parasites and Vectors, 2015, 8, 23.	2.5	60
74	Novel 2,5-disubtituted-1,3,4-oxadiazoles with benzimidazole backbone: A new class of β-glucuronidase inhibitors and in silico studies. Bioorganic and Medicinal Chemistry, 2015, 23, 3119-3125.	3.0	60
75	Identification and characterization of antibacterial compound(s) of cockroaches (Periplaneta) Tj ETQq1 1 0.784	814 rgBT /	Overlock 10 T
76	Identification and properties of proteases from an Acanthamoeba isolate capable of producing granulomatous encephalitis. BMC Microbiology, 2006, 6, 42.	3.3	59
77	Bisindolylmethane thiosemicarbazides as potential inhibitors of urease: Synthesis and molecular modeling studies. Bioorganic and Medicinal Chemistry, 2018, 26, 152-160.	3.0	59
78	Acanthamoeba is an evolutionary ancestor of macrophages: A myth or reality?. Experimental Parasitology, 2012, 130, 95-97.	1.2	57
79	The role of proteases in the differentiation of <i>Acanthamoeba castellanii</i> . FEMS Microbiology Letters, 2008, 286, 9-15.	1.8	54
80	Cytotoxic effects of aflatoxin B1 on human brain microvascular endothelial cells of the blood-brain barrier. Medical Mycology, 2015, 53, 409-416.	0.7	54
81	Silver nanoparticle conjugation affects antiacanthamoebic activities of amphotericin B, nystatin, and fluconazole. Parasitology Research, 2018, 117, 265-271.	1.6	54
82	Synthesis, biological evaluation, and docking studies of novel thiourea derivatives of bisindolylmethane as carbonic anhydrase II inhibitor. Bioorganic Chemistry, 2015, 62, 83-93.	4.1	53
83	Oxindole based oxadiazole hybrid analogs: Novel α -glucosidase inhibitors. Bioorganic Chemistry, 2018, 76, 273-280.	4.1	53
84	Oxadiazoles and thiadiazoles: Novel α-glucosidase inhibitors. Bioorganic and Medicinal Chemistry, 2014, 22, 5454-5465.	3.0	52
85	Discovery of novel oxindole derivatives as potent α-glucosidase inhibitors. Bioorganic and Medicinal Chemistry, 2014, 22, 3441-3448.	3.0	51
86	2-Arylquinazolin-4(3H)-ones: A new class of α-glucosidase inhibitors. Bioorganic and Medicinal Chemistry, 2015, 23, 7417-7421.	3.0	51
87	Combating Acanthamoeba spp. cysts: what are the options?. Parasites and Vectors, 2018, 11, 26.	2.5	51
88	Synthesis, in vitro alpha-glucosidase inhibitory potential of benzimidazole bearing bis-Schiff bases and their molecular docking study. Bioorganic Chemistry, 2020, 94, 103394.	4.1	51
89	Synthesis of new indazole based dual inhibitors of α-glucosidase and α-amylase enzymes, their in vitro, in silico and kinetics studies. Bioorganic Chemistry, 2020, 94, 103195.	4.1	51
90	<i>In Vitro</i> Efficacies of Clinically Available Drugs against Growth and Viability of an Acanthamoeba castellanii Keratitis Isolate Belonging to the T4 Genotype. Antimicrobial Agents and Chemotherapy, 2013, 57, 3561-3567.	3.2	50

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91	Biocompatible Tin Oxide Nanoparticles: Synthesis, Antibacterial, Anticandidal and Cytotoxic Activities. ChemistrySelect, 2019, 4, 4013-4017.	1.5	50
92	2-(2′-Pyridyl) benzimidazole derivatives and their urease inhibitory activity. Medicinal Chemistry Research, 2014, 23, 4447-4454.	2.4	49
93	Synthesis of Bis-indolylmethane sulfonohydrazides derivatives as potent α-Glucosidase inhibitors. Bioorganic Chemistry, 2018, 80, 112-120.	4.1	49
94	Gut microbiome and human health under the space environment. Journal of Applied Microbiology, 2021, 130, 14-24.	3.1	49
95	Synthesis of gold nanoparticles stabilized by a pyrazinium thioacetate ligand: A new colorimetric nanosensor for detection of heavy metal Pd(II). Sensors and Actuators B: Chemical, 2018, 257, 875-881.	7.8	48
96	2ʹ-Aryl and 4ʹ-arylidene substituted pyrazolones: As potential α-amylase inhibitors. European Journal of Medicinal Chemistry, 2018, 159, 47-58.	5.5	48
97	New indole based hybrid oxadiazole scaffolds with N-substituted acetamides: As potent anti-diabetic agents. Bioorganic Chemistry, 2018, 81, 253-263.	4.1	48
98	Balamuthia mandrillarisexhibits metalloprotease activities. FEMS Immunology and Medical Microbiology, 2006, 47, 83-91.	2.7	47
99	Evaluation of bisindole as potent β-glucuronidase inhibitors: Synthesis and in silico based studies. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 1825-1829.	2.2	47
100	Synthesis, α-glucosidase inhibitory, cytotoxicity and docking studies of 2-aryl-7-methylbenzimidazoles. Bioorganic Chemistry, 2016, 65, 100-109.	4.1	47
101	Dihydropyrano [2,3-c] pyrazole: Novel in vitro inhibitors of yeast α-glucosidase. Bioorganic Chemistry, 2016, 65, 61-72.	4.1	47
102	2-Aryl benzimidazoles: Synthesis, InÂvitro α-amylase inhibitory activity, and molecular docking study. European Journal of Medicinal Chemistry, 2018, 150, 248-260.	5.5	47
103	Silver Nanoparticle Conjugation-Enhanced Antibacterial Efficacy of Clinically Approved Drugs Cephradine and Vildagliptin. Antibiotics, 2018, 7, 100.	3.7	47
104	Gold Nanoparticle-Conjugated Cinnamic Acid Exhibits Antiacanthamoebic and Antibacterial Properties. Antimicrobial Agents and Chemotherapy, 2018, 62, .	3.2	47
105	Water-Pipe Smoking and Metabolic Syndrome: A Population-Based Study. PLoS ONE, 2012, 7, e39734.	2.5	46
106	Molecular modeling-based antioxidant arylidene barbiturates as urease inhibitors. Journal of Molecular Graphics and Modelling, 2011, 30, 153-156.	2.4	45
107	Pharmacological basis for the medicinal use of Linum usitatissimum (Flaxseed) in infectious and non-infectious diarrhea. Journal of Ethnopharmacology, 2015, 160, 61-68.	4.1	45
108	Synthesis of 6-chloro-2-Aryl-1H-imidazo[4,5-b]pyridine derivatives: Antidiabetic, antioxidant, β-glucuronidase inhibiton and their molecular docking studies. Bioorganic Chemistry, 2016, 65, 48-56.	4.1	45

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109	Synthesis of quinoline derivatives as diabetic II inhibitors and molecular docking studies. Bioorganic and Medicinal Chemistry, 2019, 27, 4081-4088.	3.0	45
110	Zinc oxide nanoparticles conjugated with clinically-approved medicines as potential antibacterial molecules. AMB Express, 2021, 11, 104.	3.0	45
111	Mechanisms associated with Acanthamoeba castellanii (T4) phagocytosis. Parasitology Research, 2005, 96, 402-409.	1.6	44
112	The capsule plays an important role in Escherichia coli K1 interactions with Acanthamoeba. International Journal for Parasitology, 2007, 37, 417-423.	3.1	44
113	Gut bacteria of cockroaches are a potential source of antibacterial compound(s). Letters in Applied Microbiology, 2018, 66, 416-426.	2.2	44
114	Novel Coronavirus: Current Understanding of Clinical Features, Diagnosis, Pathogenesis, and Treatment Options. Pathogens, 2020, 9, 297.	2.8	44
115	Acanthamoeba and the blood–brain barrier: the breakthrough. Journal of Medical Microbiology, 2008, 57, 1051-1057.	1.8	43
116	Synthesis and structure–activity relationship of thiobarbituric acid derivatives as potent inhibitors of urease. Bioorganic and Medicinal Chemistry, 2014, 22, 4119-4123.	3.0	43
117	Crocodiles and alligators: Antiamoebic and antitumor compounds of crocodiles. Experimental Parasitology, 2017, 183, 194-200.	1.2	43
118	Acanthamoeba castellanii: High antibody prevalence in racially and ethnically diverse populations. Experimental Parasitology, 2009, 121, 254-256.	1.2	42
119	Synthesis of novel benzohydrazone–oxadiazole hybrids as β-glucuronidase inhibitors and molecular modeling studies. Bioorganic and Medicinal Chemistry, 2015, 23, 7394-7404.	3.0	42
120	A Novel Prosumer-Based Energy Sharing and Management (PESM) Approach for Cooperative Demand Side Management (DSM) in Smart Grid. Applied Sciences (Switzerland), 2016, 6, 275.	2.5	42
121	Dihydropyridines as potential α-amylase and α-glucosidase inhibitors: Synthesis, in vitro and in silico studies. Bioorganic Chemistry, 2020, 96, 103581.	4.1	42
122	Syntheses, in vitro α-amylase and α-glucosidase dual inhibitory activities of 4-amino-1,2,4-triazole derivatives their molecular docking and kinetic studies. Bioorganic and Medicinal Chemistry, 2020, 28, 115467.	3.0	42
123	5-Acetyl-6-methyl-4-aryl-3,4-dihydropyrimidin-2(1 H)-ones: As potent urease inhibitors; synthesis, in vitro screening, and molecular modeling study. Bioorganic Chemistry, 2018, 76, 37-52.	4.1	41
124	Cobalt nanoparticles as novel nanotherapeutics against Acanthamoeba castellanii. Parasites and Vectors, 2019, 12, 280.	2.5	41
125	Indole acrylonitriles as potential anti-hyperglycemic agents: Synthesis, α-glucosidase inhibitory activity and molecular docking studies. Bioorganic and Medicinal Chemistry, 2020, 28, 115605.	3.0	41
126	Acanthamoeba isolates belonging to T1, T2, T3, T4 but not T7 encyst in response to increased osmolarity and cysts do not bind to human corneal epithelial cells. Acta Tropica, 2005, 95, 100-108.	2.0	40

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127	Expeditious Method for Synthesis of Symmetrical 1,3â€Disubstituted Ureas and Thioureas. Synthetic Communications, 2005, 35, 1663-1674.	2.1	40
128	Acanthamoeba castellanii of the T4 genotype is a potential environmental host for Enterobacter aerogenes and Aeromonas hydrophila. Parasites and Vectors, 2013, 6, 169.	2.5	40
129	Synthesis and evaluation of unsymmetrical heterocyclic thioureas as potent β-glucuronidase inhibitors. Medicinal Chemistry Research, 2015, 24, 3166-3173.	2.4	40
130	Gold Nanoparticle Conjugation Enhances the Antiacanthamoebic Effects of Chlorhexidine. Antimicrobial Agents and Chemotherapy, 2016, 60, 1283-1288.	3.2	40
131	Boron Nitride Doped Polyhydroxyalkanoate/Chitosan Nanocomposite for Antibacterial and Biological Applications. Nanomaterials, 2019, 9, 645.	4.1	40
132	Dihydropyrimidones: As novel class of β-glucuronidase inhibitors. Bioorganic and Medicinal Chemistry, 2016, 24, 3624-3635.	3.0	39
133	Chalcones and bis-chalcones: As potential α-amylase inhibitors; synthesis, in vitro screening, and molecular modelling studies. Bioorganic Chemistry, 2018, 79, 179-189.	4.1	39
134	Synthesis of azachalcones, their α-amylase, α-glucosidase inhibitory activities, kinetics, and molecular docking studies. Bioorganic Chemistry, 2021, 106, 104489.	4.1	39
135	Cellulose biosynthesis pathway is a potential target in the improved treatment of Acanthamoeba keratitis. Applied Microbiology and Biotechnology, 2007, 75, 133-140.	3.6	38
136	Protozoa traversal of the blood–brain barrier to invade the central nervous system. FEMS Microbiology Reviews, 2010, 34, 532-553.	8.6	38
137	Flurbiprofen derivatives as novel α-amylase inhibitors: Biology-oriented drug synthesis (BIODS), in vitro, and in silico evaluation. Bioorganic Chemistry, 2018, 81, 157-167.	4.1	38
138	The Use of Nanomedicine for Targeted Therapy against Bacterial Infections. Antibiotics, 2019, 8, 260.	3.7	38
139	2,4,6-Trichlorophenylhydrazine Schiff Bases as DPPH Radical and Super Oxide Anion Scavengers. Medicinal Chemistry, 2012, 8, 452-461.	1.5	38
140	Synthesis of Benzophenonehydrazone Schiff Bases and their In Vitro Antiglycating Activities. Medicinal Chemistry, 2013, 9, 588-595.	1.5	38
141	Ecto-ATPases of clinical and non-clinical isolates of Acanthamoeba. Microbial Pathogenesis, 2004, 37, 231-239.	2.9	37
142	Evaluation of 2-indolcarbohydrazones as potent α-glucosidase inhibitors, in silico studies and DFT based stereochemical predictions. Bioorganic Chemistry, 2015, 63, 24-35.	4.1	37
143	Synthesis, in vitro α-glucosidase inhibitory activity and molecular docking studies of new thiazole derivatives. Bioorganic Chemistry, 2016, 68, 245-258.	4.1	37
144	Molecular hybridization conceded exceptionally potent quinolinyl-oxadiazole hybrids through phenyl linked thiosemicarbazide antileishmanial scaffolds: In silico validation and SAR studies. Bioorganic Chemistry, 2017, 71, 192-200.	4.1	37

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145	Synthesis of piperazine sulfonamide analogs as diabetic-II inhibitors and their molecular docking study. European Journal of Medicinal Chemistry, 2017, 141, 530-537.	5.5	37
146	Inefficacy of marketed contact lens disinfection solutions against keratitis-causing Acanthamoeba castellanii belonging to the T4 genotype. Experimental Parasitology, 2014, 141, 122-128.	1.2	36
147	Acylhydrazide Schiff Bases: DPPH Radical and Superoxide Anion Scavengers. Medicinal Chemistry, 2012, 8, 705-710.	1.5	36
148	Anti-Acanthamoebic properties of resveratrol and demethoxycurcumin. Experimental Parasitology, 2012, 132, 519-523.	1.2	35
149	Dihydropyrimidine based hydrazine dihydrochloride derivatives as potent urease inhibitors. Bioorganic Chemistry, 2016, 64, 85-96.	4.1	35
150	Repositioning of Guanabenz in Conjugation with Gold and Silver Nanoparticles against Pathogenic Amoebae <i>Acanthamoeba castellanii</i> and <i>Naegleria fowleri</i> . ACS Infectious Diseases, 2019, 5, 2039-2046.	3.8	35
151	Drug Discovery against Acanthamoeba Infections: Present Knowledge and Unmet Needs. Pathogens, 2020, 9, 405.	2.8	35
152	Fabrication of biopolymer polyhydroxyalkanoate/chitosan and 2D molybdenum disulfide–doped scaffolds for antibacterial and biomedical applications. Applied Microbiology and Biotechnology, 2020, 104, 3121-3131.	3.6	35
153	Cenotypic, phenotypic, biochemical, physiological and pathogenicity-based categorisation of Acanthamoeba strains. Folia Parasitologica, 2003, 50, 97-104.	1.3	35
154	Oxindole Derivatives: Synthesis and Antiglycation Activity. Medicinal Chemistry, 2013, 9, 681-688.	1.5	35
155	Molecular Tools for Speciation and Epidemiological Studies of Acanthamoeba. Current Microbiology, 2002, 44, 444-449.	2.2	34
156	Acanthamoeba invasion of the central nervous system. International Journal for Parasitology, 2007, 37, 131-138.	3.1	34
157	War of the microbial worlds: Who is the beneficiary in Acanthamoeba–bacterial interactions?. Experimental Parasitology, 2012, 130, 311-313.	1.2	34
158	Synthesis and molecular modelling studies of phenyl linked oxadiazole-phenylhydrazone hybrids as potent antileishmanial agents. European Journal of Medicinal Chemistry, 2017, 126, 1021-1033.	5.5	34
159	Novel acridine-based thiosemicarbazones as â€~turn-on' chemosensors for selective recognition of fluoride anion: a spectroscopic and theoretical study. Royal Society Open Science, 2018, 5, 180646.	2.4	34
160	Synthesis, structure-activity relationship and molecular docking studies of 3-O-flavonol glycosides as cholinesterase inhibitors. Bioorganic and Medicinal Chemistry, 2018, 26, 3696-3706.	3.0	34
161	Synthesis, in vitro α-amylase inhibitory, and radicals (DPPH & ABTS) scavenging potentials of new N-sulfonohydrazide substituted indazoles. Bioorganic Chemistry, 2020, 94, 103410.	4.1	34
162	Antiamoebic activity of plant-based natural products and their conjugated silver nanoparticles against Acanthamoeba castellanii (ATCC 50492). AMB Express, 2020, 10, 24.	3.0	34

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