Hoon Kim

List of Publications by Year in descending order

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		117453	182168
140	3,684	34	51
papers	citations	h-index	g-index
142	142	142	2797
andocs	does citations	times ranked	citing authors

HOONKIM

#	Article	IF	CITATIONS
1	Inhibition of Monoamine Oxidase A by β-Carboline Derivatives. Archives of Biochemistry and Biophysics, 1997, 337, 137-142.	1.4	234
2	Magnetic nanoparticles for hyperthermia in cancer treatment: an emerging tool. Environmental Science and Pollution Research, 2020, 27, 19214-19225.	2.7	143
3	Advancements in nanotherapeutics for Alzheimer's disease: current perspectives. Journal of Pharmacy and Pharmacology, 2019, 71, 1370-1383.	1.2	108
4	Emerging therapeutic potentials of dualâ€acting MAO and AChE inhibitors in Alzheimer's and Parkinson's diseases. Archiv Der Pharmazie, 2019, 352, e1900177.	2.1	99
5	Cholinesterase Inhibitors for Alzheimer's Disease: Multitargeting Strategy Based on Anti-Alzheimer's Drugs Repositioning. Current Pharmaceutical Design, 2019, 25, 3519-3535.	0.9	88
6	Potent selective monoamine oxidase B inhibition by maackiain, a pterocarpan from the roots of Sophora flavescens. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 4714-4719.	1.0	82
7	Synthesis, Biochemistry, and Computational Studies of Brominated Thienyl Chalcones: A New Class of Reversible MAOâ€B Inhibitors. ChemMedChem, 2016, 11, 1161-1171.	1.6	70
8	Development of fluorinated methoxylated chalcones as selective monoamine oxidase-B inhibitors: Synthesis, biochemistry and molecular docking studies. Bioorganic Chemistry, 2015, 62, 22-29.	2.0	69
9	Exploration of chlorinated thienyl chalcones: A new class of monoamine oxidase-B inhibitors. International Journal of Biological Macromolecules, 2016, 91, 680-695.	3.6	69
10	Identification of Indoleâ€Based Chalcones: Discovery of a Potent, Selective, and Reversible Class of MAOâ€B Inhibitors. Archiv Der Pharmazie, 2016, 349, 627-637.	2.1	62
11	Molecular cloning and characterization of a novel family VIII alkaline esterase from a compost metagenomic library. Biochemical and Biophysical Research Communications, 2010, 393, 45-49.	1.0	59
12	Monoamine Oxidase Inhibitory Action of Chalcones: A Mini Review. Central Nervous System Agents in Medicinal Chemistry, 2016, 16, 120-136.	0.5	59
13	Chalcones: Unearthing their therapeutic possibility as monoamine oxidase B inhibitors. European Journal of Medicinal Chemistry, 2020, 205, 112650.	2.6	58
14	Selective inhibition of monoamine oxidase A by hispidol. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 584-588.	1.0	55
15	Heteroaryl chalcones: Mini review about their therapeutic voyage. Biomedicine and Preventive Nutrition, 2014, 4, 451-458.	0.9	52
16	Monoamine oxidase inhibitory activity of methoxy-substituted chalcones. International Journal of Biological Macromolecules, 2017, 104, 1321-1329.	3.6	51
17	Monoamine Oxidase Inhibitory Activity: Methyl―versus Chlorochalcone Derivatives. ChemMedChem, 2016, 11, 2649-2655	1.6	50
18	Potent inhibition of monoamine oxidase A by decursin from Angelica gigas Nakai and by wogonin from Scutellaria baicalensis Georgi. International Journal of Biological Macromolecules, 2017, 97, 598-605.	3.6	50

#	Article	IF	CITATIONS
19	Design, synthesis and biological evaluation of oxygenated chalcones as potent and selective MAO-B inhibitors. Bioorganic Chemistry, 2019, 93, 103335.	2.0	49
20	Rhamnocitrin isolated from Prunus padus var. seoulensis: A potent and selective reversible inhibitor of human monoamine oxidase A. Bioorganic Chemistry, 2019, 83, 317-325.	2.0	47
21	Influence of the Transposition of the Thermostabilizing Domain of Clostridium thermocellum Xylanase (XynX) on Xylan Binding and Thermostabilization. Applied and Environmental Microbiology, 2002, 68, 3496-3501.	1.4	46
22	Potent inhibition of acetylcholinesterase by sargachromanol I from Sargassum siliquastrum and by selected natural compounds. Bioorganic Chemistry, 2019, 89, 103043.	2.0	45
23	Potent and highly selective dualâ€ŧargeting monoamine oxidaseâ€B inhibitors: Fluorinated chalcones of morpholine versus imidazole. Archiv Der Pharmazie, 2019, 352, e1800309.	2.1	44
24	Pharmacophore-Based 3D-QSAR Analysis of Thienyl Chalcones as a New Class of Human MAO-B Inhibitors: Investigation of Combined Quantum Chemical and Molecular Dynamics Approach. Journal of Physical Chemistry B, 2017, 121, 1186-1203.	1.2	40
25	Selected aryl thiosemicarbazones as a new class of multi-targeted monoamine oxidase inhibitors. MedChemComm, 2018, 9, 1871-1881.	3.5	40
26	Acetylcholinesterase and butyrylcholinesterase inhibitory activities of khellactone coumarin derivatives isolated from Peucedanum japonicum Thurnberg. Scientific Reports, 2020, 10, 21695.	1.6	40
27	Potent and Selective Monoamine Oxidaseâ€B Inhibitory Activity: Fluoro― <i>vs</i> . Trifluoromethylâ€4â€hydroxylated Chalcone Derivatives. Chemistry and Biodiversity, 2016, 13, 1046-1052.	1.0	39
28	Discovery of potent and reversible MAO-B inhibitors as furanochalcones. International Journal of Biological Macromolecules, 2018, 108, 660-664.	3.6	39
29	Pyrazoline: A Promising Scaffold for the Inhibition of Monoamine Oxidase. Central Nervous System Agents in Medicinal Chemistry, 2014, 13, 195-206.	0.5	39
30	Characterization of Xyn10J, a Novel Family 10 Xylanase from a Compost Metagenomic Library. Applied Biochemistry and Biotechnology, 2012, 166, 1328-1339.	1.4	38
31	Potent inhibitions of monoamine oxidase A and B by acacetin and its 7-O-(6-O-malonylglucoside) derivative from Agastache rugosa. International Journal of Biological Macromolecules, 2017, 104, 547-553.	3.6	38
32	Development of Fluorinated Thienylchalcones as Monoamine Oxidase-B Inhibitors: Design, Synthesis, Biological Evaluation and Molecular Docking Studies. Letters in Organic Chemistry, 2015, 12, 605-613.	0.2	38
33	Imidazole bearing chalcones as a new class of monoamine oxidase inhibitors. Biomedicine and Pharmacotherapy, 2018, 106, 8-13.	2.5	36
34	Calycosin and 8-O-methylretusin isolated from Maackia amurensis as potent and selective reversible inhibitors of human monoamine oxidase-B. International Journal of Biological Macromolecules, 2020, 151, 441-448.	3.6	36
35	Novel Class of Chalcone Oxime Ethers as Potent Monoamine Oxidase-B and Acetylcholinesterase Inhibitors. Molecules, 2020, 25, 2356.	1.7	35
36	Cloning of two cellulase genes from endophytic <i>Paenibacillus polymyxa</i> GS01 and comparison with <i>cel</i> 44Câ€ <i>man</i> 26A. Journal of Basic Microbiology, 2008, 48, 464-472.	1.8	34

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37	Perspective Design of Chalcones for the Management of CNS Disorders: A Mini-Review. CNS and Neurological Disorders - Drug Targets, 2019, 18, 432-445.	0.8	34
38	Characterization of a Novel Alkaline Family VIII Esterase with S-Enantiomer Preference from a Compost Metagenomic Library. Journal of Microbiology and Biotechnology, 2016, 26, 315-325.	0.9	34
39	TV 3326 for Alzheimer's dementia: a novel multimodal ChE and MAO inhibitors to mitigate Alzheimer's-like neuropathology. Journal of Pharmacy and Pharmacology, 2020, 72, 1001-1012.	1.2	33
40	Deciphering the detailed structure–activity relationship of coumarins as Monoamine oxidase enzyme inhibitors—An updated review. Chemical Biology and Drug Design, 2021, 98, 655-673.	1,5	33
41	New Aspects of Monoamine Oxidase B Inhibitors: The Key Role of Halogens to Open the Golden Door. Current Medicinal Chemistry, 2020, 28, 266-283.	1.2	32
42	Anti-oxidant behavior of functionalized chalcone-a combined quantum chemical and crystallographic structural investigation. Journal of Molecular Structure, 2017, 1146, 301-308.	1.8	31
43	Monoamine Oxidase Inhibitors: Perspective Design for the Treatment of Depression and Neurological Disorders. Current Enzyme Inhibition, 2016, 12, 115-122.	0.3	31
44	Structural Exploration of Synthetic Chromones as Selective MAO-B Inhibitors: A Mini Review. Combinatorial Chemistry and High Throughput Screening, 2017, 20, 522-532.	0.6	30
45	Plant Secondary Metabolites- Potent Inhibitors of Monoamine Oxidase Isoforms. Central Nervous System Agents in Medicinal Chemistry, 2014, 14, 28-33.	0.5	28
46	Selective inhibition of monoamine oxidase A by chelerythrine, an isoquinoline alkaloid. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 2403-2407.	1.0	28
47	Inhibition of Butyrylcholinesterase and Human Monoamine Oxidase-B by the Coumarin Glycyrol and Liquiritigenin Isolated from Glycyrrhiza uralensis. Molecules, 2020, 25, 3896.	1.7	27
48	Ethyl Acetohydroxamate Incorporated Chalcones: Unveiling a Novel Class of Chalcones for Multitarget Monoamine Oxidase-B Inhibitors Against Alzheimer's Disease. CNS and Neurological Disorders - Drug Targets, 2019, 18, 643-654.	0.8	27
49	Activity Enhancement of Cel5Z from Pectobacterium chrysanthemi PY35 by Removing C-Terminal Region. Biochemical and Biophysical Research Communications, 2002, 291, 425-430.	1.0	26
50	Characterization of Novel Family IV Esterase and Family I.3 Lipase from an Oil-Polluted Mud Flat Metagenome. Molecular Biotechnology, 2015, 57, 781-792.	1.3	26
51	Piperazine-substituted chalcones: a new class of MAO-B, AChE, and BACE-1 inhibitors for the treatment of neurological disorders. Environmental Science and Pollution Research, 2021, 28, 38855-38866.	2.7	26
52	Halogenated Coumarin–Chalcones as Multifunctional Monoamine Oxidase-B and Butyrylcholinesterase Inhibitors. ACS Omega, 2021, 6, 28182-28193.	1.6	26
53	Selective inhibition of monoamine oxidase A by purpurin, an anthraquinone. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 1136-1140.	1.0	25
54	Potent Inhibition of Monoamine Oxidase B by a Piloquinone from Marine-Derived Streptomyces sp. CNQ-027. Journal of Microbiology and Biotechnology, 2017, 27, 785-790.	0.9	25

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55	Cloning and characterization of a novel intracellular serine protease (IspK) from Bacillus megaterium with a potential additive for detergents. International Journal of Biological Macromolecules, 2018, 108, 808-816.	3.6	24
56	Morpholine-based chalcones as dual-acting monoamine oxidase-B and acetylcholinesterase inhibitors: synthesis and biochemical investigations. Journal of Enzyme Inhibition and Medicinal Chemistry, 2021, 36, 188-197.	2.5	24
57	Development of bromo- and fluoro-based α, β-unsaturated ketones as highly potent MAO-B inhibitors for the treatment of Parkinson's disease. Journal of Molecular Structure, 2022, 1266, 133545.	1.8	24
58	Characterization of Thienylchalcones as hMAOâ€B Inhibitors: Synthesis, Biochemistry and Molecular Dynamics Studies. ChemistrySelect, 2017, 2, 11113-11119.	0.7	23
59	Characterization of a GH family 8 β-1,3-1,4-glucanase with distinctive broad substrate specificity from Paenibacillus sp. X4. Biotechnology Letters, 2015, 37, 643-655.	1.1	22
60	Unraveling the Structural Requirements of Chalcone Chemistry Towards Monoamine Oxidase Inhibition. Central Nervous System Agents in Medicinal Chemistry, 2019, 19, 6-7.	0.5	22
61	Osthenol, a prenylated coumarin, as a monoamine oxidase A inhibitor with high selectivity. Bioorganic and Medicinal Chemistry Letters, 2019, 29, 839-843.	1.0	22
62	Ethoxylated Head of Chalcones as a New Class of Multiâ€Targeted MAO Inhibitors. ChemistrySelect, 2019, 4, 6614-6619.	0.7	22
63	Trimethoxylated Halogenated Chalcones as Dual Inhibitors of MAO-B and BACE-1 for the Treatment of Neurodegenerative Disorders. Pharmaceutics, 2021, 13, 850.	2.0	22
64	Gene Therapy Approach with an Emphasis on Growth Factors: Theoretical and Clinical Outcomes in Neurodegenerative Diseases. Molecular Neurobiology, 2022, 59, 191-233.	1.9	22
65	Functional analysis of a hybrid endoglucanase of bacterial origin having a cellulose binding domain from a fungal exoglucanase. Applied Biochemistry and Biotechnology, 1998, 75, 193-204.	1.4	21
66	Pseudomonas taeanensis sp. nov., isolated from a crude oil-contaminated seashore. International Journal of Systematic and Evolutionary Microbiology, 2010, 60, 2719-2723.	0.8	20
67	Potent and selective inhibition of human monoamine oxidase-B by 4-dimethylaminochalcone and selected chalcone derivatives. International Journal of Biological Macromolecules, 2019, 137, 426-432.	3.6	19
68	Exploring the Therapeutic Potentials of Highly Selective Oxygenated Chalcone Based MAO-B Inhibitors in a Haloperidol-Induced Murine Model of Parkinson's Disease. Neurochemical Research, 2020, 45, 2786-2799.	1.6	19
69	Antimicrobial Activity of Divaricatic Acid Isolated from the Lichen Evernia mesomorpha against Methicillin-Resistant Staphylococcus aureus. Molecules, 2018, 23, 3068.	1.7	18
70	Design of enamides as new selective monoamine oxidase-B inhibitors. Journal of Pharmacy and Pharmacology, 2020, 72, 916-926.	1.2	18
71	Inhibition of monoamine oxidase A and B by demethoxycurcumin and bisdemethoxycurcumin. Journal of Applied Biological Chemistry, 2018, 61, 187-190.	0.2	18
72	Endo-β-1,4-glucanase encoded by Bacillus subtilis gene cloned in Bacillus megaterium. Enzyme and Microbial Technology, 1988, 10, 347-351.	1.6	17

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73	Characterization of two extracellular β-glucosidases produced from the cellulolytic fungus Aspergillus sp. YDJ216 and their potential applications for the hydrolysis of flavone glycosides. International Journal of Biological Macromolecules, 2018, 111, 595-603.	3.6	17
74	Selective Inhibition of Human Monoamine Oxidase B by 5-hydroxy-2-methyl-chroman-4-one Isolated from an Endogenous Lichen Fungus Daldinia fissa. Journal of Fungi (Basel, Switzerland), 2021, 7, 84.	1.5	17
75	Changes in the activity of the multifunctional β-glycosyl hydrolase (Cel44C-Man26A) from Paenibacillus polymyxa by removal of the C-terminal region to minimum size. Biotechnology Letters, 2008, 30, 1061-1068.	1.1	16
76	Characterization of truncated endo-β-1,4-glucanases from a compost metagenomic library and their saccharification potentials. International Journal of Biological Macromolecules, 2018, 115, 554-562.	3.6	15
77	Inhibition of Monoamine Oxidase by Anithiactins from Streptomyces sp Journal of Microbiology and Biotechnology, 2015, 25, 1425-1428.	0.9	15
78	Structural features of Safinamide: A combined Hirshfeld surface analysis & quantum chemical treatment. Chemical Data Collections, 2018, 17-18, 404-414.	1.1	14
79	Insights into an Immunotherapeutic Approach to Combat Multidrug Resistance in Hepatocellular Carcinoma. Pharmaceuticals, 2021, 14, 656.	1.7	14
80	Revealing the role of the benzyloxy pharmacophore in the design of a new class of monoamine oxidaseâ€B inhibitors. Archiv Der Pharmazie, 2022, 355, e2200084.	2.1	14
81	Construction of minimum size cellulase (Cel5Z) from Pectobacterium chrysanthemi PY35 by removal of the C-terminal region. Applied Microbiology and Biotechnology, 2005, 68, 46-52.	1.7	13
82	Gangjinia marincola gen. nov., sp. nov., a marine bacterium of the family Flavobacteriaceae. International Journal of Systematic and Evolutionary Microbiology, 2011, 61, 325-329.	0.8	13
83	Improvement of enzyme activity of β-1,3-1,4-glucanase from Paenibacillus sp. X4 by error-prone PCR and structural insights of mutated residues. Applied Microbiology and Biotechnology, 2017, 101, 4073-4083.	1.7	13
84	Refining the Structural Features of Chromones as Selective MAOâ€B Inhibitors: Exploration of Combined Pharmacophoreâ€Based 3Dâ€QSAR and Quantum Chemical Studies. ChemistrySelect, 2017, 2, 11645-11652.	0.7	13
85	Synthesis and biological evaluation of new 3(2H)-pyridazinone derivatives as non-toxic anti-proliferative compounds against human colon carcinoma HCT116 cells. Journal of Enzyme Inhibition and Medicinal Chemistry, 2020, 35, 1100-1109.	2.5	13
86	Synthesis, Cytotoxicity and Anti-Proliferative Activity against AGS Cells of New 3(2H)-Pyridazinone Derivatives Endowed with a Piperazinyl Linker. Pharmaceuticals, 2021, 14, 183.	1.7	13
87	Potent Selective Inhibition of Monoamine Oxidase A by Alternariol Monomethyl Ether Isolated from Alternaria brassicae. Journal of Microbiology and Biotechnology, 2017, 27, 316-320.	0.9	13
88	Biological investigation of <i>N</i> -methyl thiosemicarbazones as antimicrobial agents and bacterial carbonic anhydrases inhibitors. Journal of Enzyme Inhibition and Medicinal Chemistry, 2022, 37, 986-993.	2.5	13
89	Cloning and expression of a Clostridium thermocellum xylanase gene in Escherichia coli. IUBMB Life, 1998, 44, 283-292.	1.5	12
90	Current Progress in Quinazoline Derivatives as Acetylcholinesterase and Monoamine Oxidase Inhibitors. ChemistrySelect, 2021, 6, 7162-7182.	0.7	12

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91	Antidepressant-Like Activities of Hispidol and Decursin in Mice and Analysis of Neurotransmitter Monoamines. Neurochemical Research, 2020, 45, 1930-1940.	1.6	12
92	Privileged Pharmacophore of FDA Approved Drugs in Combination with Chalcone Framework: A New Hope for Alzheimer's Treatment. Combinatorial Chemistry and High Throughput Screening, 2020, 23, 842-846.	0.6	12
93	Selected 1,3â€Benzodioxineâ€Containing Chalcones as Multipotent Oxidase and Acetylcholinesterase Inhibitors. ChemMedChem, 2020, 15, 2257-2263.	1.6	11
94	Design, Synthesis, and Biological Evaluation of Pyridazinones Containing the (2-Fluorophenyl) Piperazine Moiety as Selective MAO-B Inhibitors. Molecules, 2020, 25, 5371.	1.7	11
95	A New Potent and Selective Monoamine Oxidaseâ€B Inhibitor with Extended Conjugation in a Chalcone Framework: 1â€[4â€(Morpholinâ€4â€yl)phenyl]â€5â€phenylpentaâ€2,4â€dienâ€1â€one. ChemMedChem, 2020	, 15 , 1629	-1633.
96	Development of methylthiosemicarbazones as new reversible monoamine oxidase-B inhibitors for the treatment of Parkinson's disease. Journal of Biomolecular Structure and Dynamics, 2021, 39, 4786-4794.	2.0	11
97	Acetylcholinesterase and monoamine oxidase-B inhibitory activities by ellagic acid derivatives isolated from Castanopsis cuspidata var. sieboldii. Scientific Reports, 2021, 11, 13953.	1.6	11
98	Navigating into the Chemical Space of Monoamine Oxidase Inhibitors by Artificial Intelligence and Cheminformatics Approach. ACS Omega, 2021, 6, 23399-23411.	1.6	11
99	Aldoxime- and hydroxy-functionalized chalcones as highly potent and selective monoamine oxidase-B inhibitors. Journal of Molecular Structure, 2022, 1250, 131817.	1.8	11
100	An Update of Synthetic Approaches and Structureâ€Activity Relationships of Various Classes of Human MAOâ€B Inhibitors. ChemistrySelect, 2021, 6, 1404-1429.	0.7	10
101	A Comprehensive Review of Monoamine Oxidase-A Inhibitors in their Syntheses and Potencies. Combinatorial Chemistry and High Throughput Screening, 2020, 23, 898-914.	0.6	10
102	Conjugated Dienones from Differently Substituted Cinnamaldehyde as Highly Potent Monoamine Oxidase-B Inhibitors: Synthesis, Biochemistry, and Computational Chemistry. ACS Omega, 2022, 7, 8184-8197.	1.6	10
103	Development of a Novel Class of Pyridazinone Derivatives as Selective MAO-B Inhibitors. Molecules, 2022, 27, 3801.	1.7	10
104	Development of Halogenated Pyrazolines as Selective Monoamine Oxidase-B Inhibitors: Deciphering via Molecular Dynamics Approach. Molecules, 2021, 26, 3264.	1.7	9
105	A cold-active acidophilic endoglucanase of <i>Paenibacillus</i> sp. Y2 isolated from soil in an alpine region. Journal of Applied Biological Chemistry, 2017, 60, 257-263.	0.2	9
106	Characterization of a Multimodular Endo-Â¥-1,4-glucanase (Cel9K) from Paenibacillus sp. X4 with a Potential Additive for Saccharification. Journal of Microbiology and Biotechnology, 2018, 28, 588-596.	0.9	9
107	Characterization of Three Extracellular �ï;½-Glucosidases Produced by a Fungal Isolate Aspergillus sp. YDJ14 and Their Hydrolyzing Activity for a Flavone Glycoside. Journal of Microbiology and Biotechnology, 2018, 28, 757-764.	0.9	9
108	Structural Modifications on Chalcone Framework for Developing New Class of Cholinesterase Inhibitors. International Journal of Molecular Sciences, 2022, 23, 3121.	1.8	9

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109	Title is missing!. Biotechnology Letters, 1997, 19, 483-486.	1.1	8
110	Enhancement of thermostability of Bacillus subtilis endoglucanase by error-prone PCR and DNA shuffling. Applied Biological Chemistry, 2017, 60, 73-78.	0.7	8
111	Chromenone Derivatives as Monoamine Oxidase Inhibitors from Marine-Derived MAR4 Clade <i>Streptomyces</i> sp. CNQ-031. Journal of Microbiology and Biotechnology, 2021, 31, 1022-1027.	0.9	8
112	Ameliorative effect of ethoxylated chalcone-based MAO-B inhibitor on behavioural predictors of haloperidol-induced Parkinsonism in mice: evidence of its antioxidative role against Parkinson's diseases. Environmental Science and Pollution Research, 2022, 29, 7271-7282.	2.7	8
113	Chitinibacter suncheonensis sp. nov., a chitinolytic bacterium from a mud flat in Suncheon Bay. Journal of Microbiology, 2012, 50, 1058-1062.	1.3	7
114	Enhanced saccharification of reed and rice straws by the addition of β-1,3-1,4-glucanase with broad substrate specificity and calcium ion. Journal of the Korean Society for Applied Biological Chemistry, 2015, 58, 29-33.	0.9	7
115	Antidepressant-Like Effects of Ethanol Extract of Ziziphus jujuba Mill Seeds in Mice. Applied Sciences (Switzerland), 2020, 10, 7374.	1.3	7
116	Molecular Characterization of Novel Family IV and VIII Esterases from a Compost Metagenomic Library. Microorganisms, 2021, 9, 1614.	1.6	7
117	Selected Class of Enamides Bearing Nitro Functionality as Dual-Acting with Highly Selective Monoamine Oxidase-B and BACE1 Inhibitors. Molecules, 2021, 26, 6004.	1.7	7
118	Biochemical characterization of a family IV esterase with R-form enantioselectivity from a compost metagenomic library. Applied Biological Chemistry, 2021, 64, .	0.7	7
119	Replacement of Chalcone-Ethers with Chalcone-Thioethers as Potent and Highly Selective Monoamine Oxidase-B Inhibitors and Their Protein-Ligand Interactions. Pharmaceuticals, 2021, 14, 1148.	1.7	7
120	Cleavage of bacillus subtilis endo-?-1,4-glucanase by B. megaterium protease. Biotechnology Letters, 1993, 15, 127-132.	1.1	6
121	Discovery of some novel imines of 2-amino, 5-thio, 1,3,4-thiadiazole as mucomembranous protector. Synthesis, anti-oxidant activity and in silico PASS approach. Journal of Saudi Chemical Society, 2016, 20, S426-S432.	2.4	6
122	Characterization of an Alkaline Family I.4 Lipase from Bacillus sp. W130-35 Isolated from a Tidal Mud Flat with Broad Substrate Specificity. Journal of Microbiology and Biotechnology, 2015, 25, 2024-2033.	0.9	6
123	Potent and Selective Inhibitors of Human Monoamine Oxidase A from an Endogenous Lichen Fungus Diaporthe mahothocarpus. Journal of Fungi (Basel, Switzerland), 2021, 7, 876.	1.5	6
124	(Hetero-)(arylidene)arylhydrazides as Multitarget-Directed Monoamine Oxidase Inhibitors. ACS Combinatorial Science, 2020, 22, 592-599.	3.8	5
125	Revealing the role of fluorine pharmacophore in chalcone scaffold for shifting the MAO-B selectivity: investigation of a detailed molecular dynamics and quantum chemical study. Journal of Biomolecular Structure and Dynamics, 2021, 39, 6126-6139.	2.0	5
126	Characterization of a Novel Family IV Esterase Containing a Predicted CzcO Domain and a Family V Esterase with Broad Substrate Specificity from an Oil-Polluted Mud Flat Metagenomic Library. Applied Sciences (Switzerland), 2021, 11, 5905.	1.3	5

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127	Inhibitions of monoamine oxidases and acetylcholinesterase by 1-methyl, 5-phenyl substituted thiosemicarbazones: Synthesis, biochemical, and computational investigations. Process Biochemistry, 2020, 99, 246-253.	1.8	5
128	Inhibitors of Monoamine Oxidase and Acetylcholinesterase as a Front Runner in CNS Drug Discovery. Combinatorial Chemistry and High Throughput Screening, 2020, 23, 834-835.	0.6	5
129	(S)-5-Methylmellein Isolated from an Endogenous Lichen Fungus Rosellinia corticium as a Potent Inhibitor of Human Monoamine Oxidase A. Processes, 2022, 10, 166.	1.3	5
130	Analysis of Active Metabolites of Sophora flavescens for Indoleamine 2,3-dioxygenase and Monoamine Oxidases using Ultra-Performance Liquid Chromatography-Quadrupole time-of-Flight Mass Spectrometry. Natural Product Communications, 2018, 13, 1934578X1801301.	0.2	4
131	Synthesis of N′-(4-/3-/2-/Non-substituted benzylidene)-4-[(4-methylphenyl)sulfonyloxy] Benzohydrazides and Evaluation of Their Inhibitory Activities against Monoamine Oxidases and β-Secretase. Applied Sciences (Switzerland), 2021, 11, 5830.	1.3	4
132	An Environment-friendly Synthesis of Piperonal Chalcones and Their Cytotoxic and Antioxidant Evaluation. Letters in Drug Design and Discovery, 2020, 17, 138-144.	0.4	4
133	Evaluation of Inhibitory Activities of Sophora flavescens and Angelica gigas Nakai Root Extracts against Monoamine Oxidases, Cholinesterases, and β-Secretase. Processes, 2022, 10, 880.	1.3	4
134	Title is missing!. Biotechnology Letters, 1997, 19, 27-29.	1.1	3
135	Characterization of a metalloprotease from an isolate Bacillus thuringiensis 29-126 in animal feces collected from a zoological garden in Japan. Journal of Applied Biological Chemistry, 2016, 59, 373-377.	0.2	3
136	Synthesis of New 1-Aryl-2-(3,5-dimethylpyrazol-1-yl)ethanone Oxime Ether Derivatives and Investigation of Their Cytotoxic Effects. Processes, 2021, 9, 2019.	1.3	2
137	Roles of Carbohydrate-Binding Module (CBM) of an Endo-β-1,4-Glucanase (Cel5L) from sp. KD1014 in Thermostability and Small-Substrate Hydrolyzing Activity. Journal of Microbiology and Biotechnology, 2018, 28, 2036-2045.	0.9	2
138	Adsorption of Bacillus subtilis endo-l²-1,4-glucanase to cellulosic materials. IUBMB Life, 1997, 41, 665-677.	1.5	1
139	Characterization of an alkaline esterase from an enriched metagenomic library derived from an oil-spill area. Journal of Applied Biological Chemistry, 2019, 62, 73-79.	0.2	1
140	Molecular Characterization of Four Alkaline Chitinases from Three Chitinolytic Bacteria Isolated from a Mudflat. International Journal of Molecular Sciences, 2021, 22, 12822.	1.8	1