

# Mahmud Tareq Hassan Khan

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

32 papers	1,853 citations	22 h-index	32 g-index
32 ext. papers	2,120 ext. citations	3.9 avg, IF	4.86 L-index

#	Paper	IF	Citations
32	A comprehensive review on tyrosinase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2019</b> , 34, 279-309	5.6	331
31	Retrained Classification of Tyrosinase Inhibitors and In Silico Potency Estimation by Using Atom-Type Linear Indices. <i>International Journal of Chemoinformatics and Chemical Engineering</i> , <b>2012</b> , 2, 42-144	0.3	
30	Dipeptide inhibitors of thermolysin and angiotensin I-converting enzyme. <i>Current Topics in Medicinal Chemistry</i> , <b>2012</b> , 12, 1748-62	3	3
29	Ligand-based computer-aided discovery of tyrosinase inhibitors. Applications of the TOMOCOMD-CARDD method to the elucidation of new compounds. <i>Current Pharmaceutical Design</i> , <b>2010</b> , 16, 2601-24	3.3	33
28	Predictions of the ADMET properties of candidate drug molecules utilizing different QSAR/QSPR modelling approaches. <i>Current Drug Metabolism</i> , <b>2010</b> , 11, 285-95	3.5	107
27	Tyrosinase inhibitory effect of benzoic acid derivatives and their structure-activity relationships. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2010</b> , 25, 812-7	5.6	8
26	Identification of novel quinazolin-4(3H)-ones as inhibitors of thermolysin, the prototype of the M4 family of proteinases. <i>Bioorganic and Medicinal Chemistry</i> , <b>2010</b> , 18, 4317-27	3.4	21
25	Cholinesterase inhibitory activities of some flavonoid derivatives and chosen xanthone and their molecular docking studies. <i>Chemico-Biological Interactions</i> , <b>2009</b> , 181, 383-9	5	156
24	Recent advancements for the evaluation of anti-viral activities of natural products. <i>New Biotechnology</i> , <b>2009</b> , 25, 347-68	6.4	71
23	Molecular interactions of cholinesterases inhibitors using in silico methods: current status and future prospects. <i>New Biotechnology</i> , <b>2009</b> , 25, 331-46	6.4	48
22	Discovery of potent thermolysin inhibitors using structure based virtual screening and binding assays. <i>Journal of Medicinal Chemistry</i> , <b>2009</b> , 52, 48-61	8.3	26
21	Determinants for psychrophilic and thermophilic features of metallopeptidases of the M4 family. <i>In Silico Biology</i> , <b>2009</b> , 9, 105-24	2	2
20	Ethnomedicines and ethnomedicinal phytophores against herpesviruses. <i>Biotechnology Annual Review</i> , <b>2008</b> , 14, 297-348		26
19	Atom- and bond-based 2D TOMOCOMD-CARDD approach and ligand-based virtual screening for the drug discovery of new tyrosinase inhibitors. <i>Journal of Biomolecular Screening</i> , <b>2008</b> , 13, 1014-24		23
18	Potentials of phenolic molecules of natural origin and their derivatives as anti-HIV agents. <i>Biotechnology Annual Review</i> , <b>2007</b> , 13, 223-64		18
17	Prediction of tyrosinase inhibition activity using atom-based bilinear indices. <i>ChemMedChem</i> , <b>2007</b> , 2, 449-78	3.7	43
16	TOMOCOMD-CARDD descriptors-based virtual screening of tyrosinase inhibitors: evaluation of different classification model combinations using bond-based linear indices. <i>Bioorganic and Medicinal Chemistry</i> , <b>2007</b> , 15, 1483-503	3.4	68

15	Dragon method for finding novel tyrosinase inhibitors: Biosilico identification and experimental in vitro assays. <i>European Journal of Medicinal Chemistry</i> , <b>2007</b> , 42, 1370-81	6.8	57
14	Bond-based 2D TOMOCOMD-CARDD approach for drug discovery: aiding decision-making in 'in silico' selection of new lead tyrosinase inhibitors. <i>Journal of Computer-Aided Molecular Design</i> , <b>2007</b> , 21, 167-88	4.2	18
13	Predictive QSAR modeling for the successful predictions of the ADMET properties of candidate drug molecules. <i>Current Drug Discovery Technologies</i> , <b>2007</b> , 4, 141-9	1.5	26
12	Molecular design of tyrosinase inhibitors: A critical review of promising novel inhibitors from synthetic origins. <i>Pure and Applied Chemistry</i> , <b>2007</b> , 79, 2277-2295	2.1	70
11	Heterocyclic Compounds Against the Enzyme Tyrosinase Essential for Melanin Production: Biochemical Features of Inhibition <b>2007</b> , 119-138		11
10	New tyrosinase inhibitors selected by atomic linear indices-based classification models. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2006</b> , 16, 324-30	2.9	48
9	Tyrosinase inhibition studies of cycloartane and cucurbitane glycosides and their structure-activity relationships. <i>Bioorganic and Medicinal Chemistry</i> , <b>2006</b> , 14, 6085-8	3.4	24
8	Tetraketones: a new class of tyrosinase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , <b>2006</b> , 14, 344-51	3.4	83
7	Oxazolones: new tyrosinase inhibitors; synthesis and their structure-activity relationships. <i>Bioorganic and Medicinal Chemistry</i> , <b>2006</b> , 14, 6027-33	3.4	79
6	Microbial transformation of 17alpha-ethynyl- and 17alpha-ethylsteroids, and tyrosinase inhibitory activity of transformed products. <i>Steroids</i> , <b>2005</b> , 70, 798-802	2.8	30
5	Expression of Estrogen Receptor $\alpha$ Gene in Breast Cancer Cells Treated With Transcription Factor Decoy Is Modulated by Bangladeshi Natural Plant Extracts. <i>Oncology Research</i> , <b>2005</b> , 15, 69-79	4.8	21
4	Structure-activity relationships of tyrosinase inhibitory combinatorial library of 2,5-disubstituted-1,3,4-oxadiazole analogues. <i>Bioorganic and Medicinal Chemistry</i> , <b>2005</b> , 13, 3385-95	3.4	144
3	Extracts and molecules from medicinal plants against herpes simplex viruses. <i>Antiviral Research</i> , <b>2005</b> , 67, 107-19	10.8	200
2	Synthesis of methyl ether analogues of sildenafil (Viagra) possessing tyrosinase inhibitory potential. <i>Chemistry and Biodiversity</i> , <b>2005</b> , 2, 470-6	2.5	17
1	Tyrosinase inhibitors from <i>Rhododendron collettianum</i> and their structure-activity relationship (SAR) studies. <i>Chemical and Pharmaceutical Bulletin</i> , <b>2004</b> , 52, 1458-61	1.9	41