List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Synthesis and antimicrobial activity of novel 2-thiazolylimino-5-arylidene-4-thiazolidinones. Bioorganic and Medicinal Chemistry, 2006, 14, 3859-3864.	1.4	272
2	Novel 4-thiazolidinone derivatives as potential antifungal and antibacterial drugs. Bioorganic and Medicinal Chemistry, 2010, 18, 426-432.	1.4	220
3	Antioxidants and Inflammatory Disease: Synthetic and Natural Antioxidants with Anti-Inflammatory Activity. Combinatorial Chemistry and High Throughput Screening, 2006, 9, 425-442.	0.6	196
4	Computer-Aided Discovery of Anti-Inflammatory Thiazolidinones with Dual Cyclooxygenase/Lipoxygenase Inhibition. Journal of Medicinal Chemistry, 2008, 51, 1601-1609.	2.9	161
5	2-Heteroarylimino-5-benzylidene-4-thiazolidinones analogues of 2-thiazolylimino-5-benzylidene-4-thiazolidinones with antimicrobial activity: Synthesis and structure–activity relationship. Bioorganic and Medicinal Chemistry, 2008, 16, 3714-3724.	1.4	138
6	Thiazole-based chalcones as potent antimicrobial agents. Synthesis and biological evaluation. Bioorganic and Medicinal Chemistry, 2011, 19, 3135-3140.	1.4	128
7	Design, synthesis, computational and biological evaluation of new anxiolytics. Bioorganic and Medicinal Chemistry, 2004, 12, 6559-6568.	1.4	114
8	Thiazole Ring—A Biologically Active Scaffold. Molecules, 2021, 26, 3166.	1.7	114
9	Thiazoles and Thiazolidinones as COX/LOX Inhibitors. Molecules, 2018, 23, 685.	1.7	110
10	Adamantane derivatives of thiazolyl-N-substituted amide, as possible non-steroidal anti-inflammatory agents. European Journal of Medicinal Chemistry, 2009, 44, 1198-1204.	2.6	100
11	2-Thiazolylimino/Heteroarylimino-5-arylidene-4-thiazolidinones as New Agents with SHP-2 Inhibitory Action. Journal of Medicinal Chemistry, 2008, 51, 5221-5228.	2.9	98
12	Synthesis of some new S-triazine based chalcones and their derivatives as potent antimicrobial agents. European Journal of Medicinal Chemistry, 2010, 45, 510-518.	2.6	92
13	In vitro antioxidant activity of thiazolidinone derivatives of 1,3-thiazole and 1,3,4-thiadiazole. Chemico-Biological Interactions, 2018, 286, 119-131.	1.7	81
14	Design of New Cognition Enhancers:  From Computer Prediction to Synthesis and Biological Evaluation. Journal of Medicinal Chemistry, 2004, 47, 2870-2876.	2.9	75
15	Fragment-based design, docking, synthesis, biological evaluation and structure–activity relationships of 2-benzo/benzisothiazolimino-5-aryliden-4-thiazolidinones as cycloxygenase/lipoxygenase inhibitors. European Journal of Medicinal Chemistry, 2012, 47, 111-124.	2.6	72
16	Synthesis, physicochemical characterization, cytotoxicity, antimicrobial, anti-inflammatory and psychotropic activity of new N-[1,3-(benzo)thiazol-2-yl]-1‰-[3,4-dihydroisoquinolin-2(1H)-yl]alkanamides. European Journal of Medicinal Chemistry, 2013, 70, 846-856.	2.6	63
17	Synthesis and biological evaluation of some 5-arylidene-2-(1,3-thiazol-2-ylimino)-1,3-thiazolidin-4-ones as dual anti-inflammatory/antimicrobial agents. Bioorganic and Medicinal Chemistry, 2013, 21, 532-539.	1.4	61
18	Sulfonamide-1,2,4-thiadiazole Derivatives as Antifungal and Antibacterial Agents: Synthesis, Biological Evaluation, Lipophilicity, and Conformational Studies. Chemical and Pharmaceutical Bulletin, 2010, 58, 160-167.	0.6	60

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#	Article	IF	CITATIONS
19	Thiazoles and Thiazolidinones as Antioxidants. Current Medicinal Chemistry, 2013, 20, 4460-4480.	1.2	60
20	5-Adamantan thiadiazole-based thiazolidinones as antimicrobial agents. Design, synthesis, molecular docking and evaluation. Bioorganic and Medicinal Chemistry, 2018, 26, 4664-4676.	1.4	57
21	In Silico Evaluation of the Effectivity of Approved Protease Inhibitors against the Main Protease of the Novel SARS-CoV-2 Virus. Molecules, 2020, 25, 2529.	1.7	55
22	New Benzothiazole-based Thiazolidinones as Potent Antimicrobial Agents. Design, synthesis and Biological Evaluation. Current Topics in Medicinal Chemistry, 2018, 18, 75-87.	1.0	51
23	PTP1b Inhibition, A Promising Approach for the Treatment of Diabetes Type II. Current Topics in Medicinal Chemistry, 2019, 19, 246-263.	1.0	49
24	Computer-aided prediction for medicinal chemistry via the Internet. SAR and QSAR in Environmental Research, 2008, 19, 27-38.	1.0	44
25	Evaluation of the local anaesthetic activity of 3-aminobenzo[d]isothiazole derivatives using the rat sciatic nerve model. European Journal of Medicinal Chemistry, 2009, 44, 473-481.	2.6	43
26	4-Thiazolidinone derivatives as potent antimicrobial agents: microwave-assisted synthesis, biological evaluation and docking studies. MedChemComm, 2015, 6, 319-326.	3.5	41
27	Design, synthesis and antimicrobial activity of usnic acid derivatives. MedChemComm, 2018, 9, 870-882.	3.5	40
28	Aldose reductase and protein tyrosine phosphatase 1B inhibitors as a promising therapeutic approach for diabetes mellitus. European Journal of Medicinal Chemistry, 2020, 207, 112742.	2.6	36
29	Title is missing!. Chemistry of Heterocyclic Compounds, 2002, 38, 859-866.	0.6	35
30	Novel thiazolyl, thiazolinyl and benzothiazolyl Schiff bases as possible lipoxygenase's inhibitors and anti-inflammatory agents. Il Farmaco, 2003, 58, 489-495.	0.9	35
31	Application of Docking Analysis in the Prediction and Biological Evaluation of the Lipoxygenase Inhibitory Action of Thiazolyl Derivatives of Mycophenolic Acid. Molecules, 2018, 23, 1621.	1.7	30
32	Thiazole-Based Thiazolidinones as Potent Antimicrobial Agents. Design, Synthesis and Biological Evaluation. Combinatorial Chemistry and High Throughput Screening, 2016, 19, 51-57.	0.6	29
33	New vinyl-1,2,4-triazole derivatives as antimicrobial agents: Synthesis, biological evaluation and molecular docking studies. Bioorganic and Medicinal Chemistry Letters, 2020, 30, 127368.	1.0	29
34	Synthesis and structure of condensed triazolo- and tetrazolopyrimidines. Tetrahedron, 2013, 69, 10637-10643.	1.0	28
35	Heteroarylimino-4-thiazolidinones as inhibitors of cartilage degradation. Bioorganic Chemistry, 2011, 39, 48-52.	2.0	27
36	Potent, orally available, selective COX-2 inhibitors based on 2-imidazoline core. European Journal of Medicinal Chemistry, 2014, 84, 160-172.	2.6	27

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37	Chromenone derivatives as a versatile scaffold with dual mode of inhibition of HIV-1 reverse transcriptase-associated Ribonuclease H function and integrase activity. European Journal of Medicinal Chemistry, 2019, 182, 111617.	2.6	27
38	Aminothiazole derivatives with antidegenerative activity on cartilage. Bioorganic and Medicinal Chemistry, 2003, 11, 2983-2989.	1.4	26
39	Synthesis and Biological Evaluation of Potent Antifungal Agents. Current Topics in Medicinal Chemistry, 2013, 13, 2684-2733.	1.0	25
40	Design, synthesis and biological evaluation of new substituted 5-benzylideno-2-adamantylthiazol[3,2-b][1,2,4]triazol-6(5 H)ones. Pharmacophore models for antifungal activity. Arabian Journal of Chemistry, 2018, 11, 573-590.	2.3	25
41	Heterocycle Compounds with Antimicrobial Activity. Current Pharmaceutical Design, 2020, 26, 867-904.	0.9	25
42	Novel Thiazolidin-4-ones as Potential Non-nucleoside Inhibitors of HIV-1 Reverse Transcriptase. Molecules, 2019, 24, 3821.	1.7	24
43	Design, Synthesis, Evaluation of Antimicrobial Activity and Docking Studies of New Thiazole-based Chalcones. Current Topics in Medicinal Chemistry, 2019, 19, 356-375.	1.0	23
44	Thiazolidin-4-Ones as Potential Antimicrobial Agents: Experimental and In Silico Evaluation. Molecules, 2022, 27, 1930.	1.7	23
45	Synthesis and HIV-1 RT inhibitory action of novel (4/6-substituted benzo[d]thiazol) Tj ETQq1 1 0.784314 rgBT /Ov Inhibition and Medicinal Chemistry, 2013, 28, 113-122.	verlock 10 2.5	Tf 50 427 T 22
46	Discovery of novel JAK2 and EGFR inhibitors from a series of thiazole-based chalcone derivatives. RSC Medicinal Chemistry, 2021, 12, 430-438.	1.7	22
46 47	Discovery of novel JAK2 and EGFR inhibitors from a series of thiazole-based chalcone derivatives. RSC Medicinal Chemistry, 2021, 12, 430-438. Inhibition of carbonic anhydrase isoforms I, II, IX and XII with secondary sulfonamides incorporating benzothiazole scaffolds. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1306-1311.	1.7 2.5	22 20
46 47 48	Discovery of novel JAK2 and EGFR inhibitors from a series of thiazole-based chalcone derivatives. RSC Medicinal Chemistry, 2021, 12, 430-438. Inhibition of carbonic anhydrase isoforms I, II, IX and XII with secondary sulfonamides incorporating benzothiazole scaffolds. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1306-1311. N-Heterocyclic choline analogues based on 1,2,3,4-tetrahydro(iso)quinoline scaffold with anticancer and anti-infective dual action. Pharmacological Reports, 2017, 69, 575-581.	1.7 2.5 1.5	22 20 20
46 47 48 49	Discovery of novel JAK2 and EGFR inhibitors from a series of thiazole-based chalcone derivatives. RSC Medicinal Chemistry, 2021, 12, 430-438. Inhibition of carbonic anhydrase isoforms I, II, IX and XII with secondary sulfonamides incorporating benzothiazole scaffolds. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1306-1311. N-Heterocyclic choline analogues based on 1,2,3,4-tetrahydro(iso)quinoline scaffold with anticancer and anti-infective dual action. Pharmacological Reports, 2017, 69, 575-581. Synthesis, Biological Evaluation, and Molecular Docking Studies. Molecules, 2020, 25, 1964.	1.7 2.5 1.5 1.7	22 20 20 20
46 47 48 49 50	Discovery of novel JAK2 and EGFR inhibitors from a series of thiazole-based chalcone derivatives. RSC Medicinal Chemistry, 2021, 12, 430-438. Inhibition of carbonic anhydrase isoforms I, II, IX and XII with secondary sulfonamides incorporating benzothiazole scaffolds. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1306-1311. N-Heterocyclic choline analogues based on 1,2,3,4-tetrahydro(iso)quinoline scaffold with anticancer and anti-infective dual action. Pharmacological Reports, 2017, 69, 575-581. Synthesis, Biological Evaluation, and Molecular Docking Studies. Molecules, 2020, 25, 1964. Computer Aided Predicting the Biological Activity Spectra and Experimental Testing of New Thiazole Derivatives. QSAR and Combinatorial Science, 1999, 18, 16-25.	1.7 2.5 1.5 1.7 1.4	22 20 20 20 19
 46 47 48 49 50 51 	Discovery of novel JAK2 and EGFR inhibitors from a series of thiazole-based chalcone derivatives. RSC Medicinal Chemistry, 2021, 12, 430-438. Inhibition of carbonic anhydrase isoforms I, II, IX and XII with secondary sulfonamides incorporating benzothiazole scaffolds. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1306-1311. N-Heterocyclic choline analogues based on 1,2,3,4-tetrahydro(iso)quinoline scaffold with anticancer and anti-infective dual action. Pharmacological Reports, 2017, 69, 575-581. Synthesis, Biological Evaluation, and Molecular Docking Studies. Molecules, 2020, 25, 1964. Computer Aided Predicting the Biological Activity Spectra and Experimental Testing of New Thiazole Derivatives. QSAR and Combinatorial Science, 1999, 18, 16-25. Antibacterial activity of griseofulvin analogues as an example of drug repurposing. International Journal of Antimicrobial Agents, 2020, 55, 105884.	 1.7 2.5 1.5 1.7 1.4 1.1 	22 20 20 20 19
 46 47 48 49 50 51 52 	Discovery of novel JAK2 and EGFR inhibitors from a series of thiazole-based chalcone derivatives. RSC Medicinal Chemistry, 2021, 12, 430-438. Inhibition of carbonic anhydrase isoforms I, II, IX and XII with secondary sulfonamides incorporating benzothiazole scaffolds. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1306-1311. N-Heterocyclic choline analogues based on 1,2,3,4-tetrahydro(iso)quinoline scaffold with anticancer and anti-Infective dual action. Pharmacological Reports, 2017, 69, 575-581. Synthesis, Biological Evaluation, and Molecular Docking Studies. Molecules, 2020, 25, 1964. Computer Aided Predicting the Biological Activity Spectra and Experimental Testing of New Thiazole Derivatives. QSAR and Combinatorial Science, 1999, 18, 16-25. Antibacterial activity of griseoful/in analogues as an example of drug repurposing. International Journal of Antimicrobial Agents, 2020, 55, 105884. Novel (E)-1-(4-methyl-2-(alkylamino)thiazol-5-yl)-3-arylprop-2-en-1-ones as potent antimicrobial agents. Bioorganic and Medicinal Chemistry, 2011, 19, 7349-7356.	 1.7 2.5 1.5 1.7 1.4 1.4 1.4 	22 20 20 20 19 19 18
 46 47 48 49 50 51 52 53 	Discovery of novel JAK2 and EGFR inhibitors from a series of thiazole-based chalcone derivatives. RSC Medicinal Chemistry, 2021, 12, 430-438. Inhibition of carbonic anhydrase isoforms I, II, IX and XII with secondary sulfonamides incorporating benzothiazole scaffolds. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1306-1311. N-Heterocyclic choline analogues based on 1,2,3,4-tetrahydro(iso)quinoline scaffold with anticancer and anti-infective dual action. Pharmacological Reports, 2017, 69, 575-581. Synthesis, Biological Evaluation, and Molecular Docking Studies. Molecules, 2020, 25, 1964. Computer Aided Predicting the Biological Activity Spectra and Experimental Testing of New Thiazole Derivatives. QSAR and Combinatorial Science, 1999, 18, 16-25. Antibacterial activity of griseofulvin analogues as an example of drug repurposing. International Journal of Antimicrobial Agents, 2020, 55, 105884. Novel (E)-1-(4-methyl-2-(alkylamino)thiazol-5-yl)-3-arylprop-2-en-1-ones as potent antimicrobial agents. Bioorganic and Medicinal Chemistry, 2011, 19, 7349-7356. Synthesis, Antitumor Activity, and Docking Analysis of New Pyrido[3候,2候:4,5]furo(thieno)[3,2-d]pyrimidin-8-amines. Molecules, 2019, 24, 3952.	 1.7 2.5 1.5 1.7 1.4 1.4 1.4 1.7 	22 20 20 20 19 19 18

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55	2-Aryl-3-(6-trifluoromethoxy)benzo[d]thiazole-based thiazolidinone hybrids as potential anti-infective agents: Synthesis, biological evaluation and molecular docking studies. Bioorganic and Medicinal Chemistry Letters, 2021, 32, 127718.	1.0	18
56	New Caffeic Acid Derivatives as Antimicrobial Agents: Design, Synthesis, Evaluation and Docking. Current Topics in Medicinal Chemistry, 2019, 19, 292-304.	1.0	18
57	Computer aided prediction of biological activity spectra: Evaluating versus known and predicting of new activities for thiazole derivatives. SAR and QSAR in Environmental Research, 2002, 13, 457-471.	1.0	17
58	New heterocyclic systems derived from pyridine: new substrates forÂthe investigation of the azide/tetrazole equilibrium. Tetrahedron, 2014, 70, 8648-8656.	1.0	17
59	Rational Use of Heterogeneous Data in Quantitative Structure–Activity Relationship (QSAR) Modeling of Cyclooxygenase/Lipoxygenase Inhibitors. Journal of Chemical Information and Modeling, 2019, 59, 713-730.	2.5	17
60	Appendix A. dithioloquinolinethiones as new potential multitargeted antibacterial and antifungal agents: Synthesis, biological evaluation and molecular docking studies. European Journal of Medicinal Chemistry, 2019, 175, 201-214.	2.6	17
61	5-Benzyliden-2-(5-methylthiazol-2-ylimino)thiazolidin-4-ones as Antimicrobial Agents. Design, Synthesis, Biological Evaluation and Molecular Docking Studies. Antibiotics, 2021, 10, 309.	1.5	17
62	Triazolo Based-Thiadiazole Derivatives. Synthesis, Biological Evaluation and Molecular Docking Studies. Antibiotics, 2021, 10, 804.	1.5	17
63	Prediction of enzyme inhibition and mode of inhibitory action based on calculation of distances between hydrogen bond donor/acceptor groups of the molecule and docking analysis: An application on the discovery of novel effective PTP1B inhibitors. SAR and QSAR in Environmental Research, 2015, 26.557-576	1.0	16
64	3-Amino-5-(indol-3-yl)methylene-4-oxo-2-thioxothiazolidine Derivatives as Antimicrobial Agents: Synthesis, Computational and Biological Evaluation. Pharmaceuticals, 2020, 13, 229.	1.7	16
65	Thiazole-based Chalcone Derivatives as Potential Anti-inflammatory Agents: Biological Evaluation and Molecular Modelling. Current Topics in Medicinal Chemistry, 2021, 21, 257-268.	1.0	16
66	New Substituted 5-Benzylideno-2-Adamantylthiazol[3,2-b][1,2,4]Triazol-6(5H)ones as Possible Anti-Inflammatory Agents. Molecules, 2021, 26, 659.	1.7	16
67	Thiazole/Thiadiazole/Benzothiazole Based Thiazolidin-4-One Derivatives as Potential Inhibitors of Main Protease of SARS-CoV-2. Molecules, 2022, 27, 2180.	1.7	16
68	On the reactivity of pyrido[3′,2′:4,5]furo(thieno)[3,2-d]pyrimidin-7(8)-ones with some alkyl mono- and di-halides: synthesis of new heterocyclic systems containing thiazolo[3,2-a]pyrimidine and pyrimido[2,1-b]thiazine moiety. Tetrahedron, 2015, 71, 7638-7646.	1.0	14
69	Docking assisted design of novel 4-adamantanyl-2-thiazolylimino-5-arylidene-4-thiazolidinones as potent NSAIDs. SAR and QSAR in Environmental Research, 2018, 29, 83-101.	1.0	14
70	Recent Trends in Enzyme Inhibition and Activation in Drug Design. Molecules, 2021, 26, 17.	1.7	14
71	Pyridofuropyrrolo[1,2-a]pyrimidines and pyridofuropyrimido[1,2-a]azepines: new chemical entities (NCE) with anticonvulsive and psychotropic properties. RSC Advances, 2016, 6, 49028-49038.	1.7	13
72	Synthesis and antimicrobial activity of new derivatives of pyrano[4'',3'':4',5']pyrido[3',2':4,5]thieno[3,2- <i>d</i>)pyrimidine and new heterocyclic systems. Synthetic Communications, 2019, 49, 1262-1276.	1.1	13

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73	Novel Thiazolidinone Derivatives with an Uncommon Mechanism of Inhibition Towards HIV-1 Reverse Transcriptase. Letters in Drug Design and Discovery, 2010, 7, 228-234.	0.4	13
74	Thiazolyl-N-substituted amides: A group of effective anti-inflammatory agents with potential for local anesthetic properties. Synthesis, biological evaluation, and a QSAR approach. Drug Development Research, 1999, 48, 53-60.	1.4	12
75	Thiazole-based aminopyrimidines and N-phenylpyrazolines as potent antimicrobial agents: synthesis and biological evaluation. MedChemComm, 2014, 5, 915-922.	3.5	12
76	Synthesis and antimicrobial activity of new amino derivatives of pyrano[4'',3'':4',5']pyrido[3',2':4,5]thieno[3,2-d]pyrimidine. Anais Da Academia 90, 1043-1057.	Br o stleira	De1©iencias, 2
77	Antimicrobial Activity of Nitrogen-Containing 5-α-Androstane Derivatives: In Silico and Experimental Studies. Antibiotics, 2020, 9, 224.	1.5	12
78	Non-acidic bifunctional benzothiazole-based thiazolidinones with antimicrobial and aldose reductase inhibitory activity as a promising therapeutic strategy for sepsis. Medicinal Chemistry Research, 2021, 30, 1837-1848.	1.1	12
79	Griseofulvin Derivatives: Synthesis, Molecular Docking and Biological Evaluation. Current Topics in Medicinal Chemistry, 2019, 19, 1145-1161.	1.0	12
80	Organosilicon-Containing Thiazole Derivatives as Potential Lipoxygenase Inhibitors and Anti-Inflammatory Agents. Bioinorganic Chemistry and Applications, 2007, 2007, 1-7.	1.8	11
81	On the reaction of 2-[(4-cyano-5,6,7,8-tetrahydroisoquinolin-3-yl)oxy]acetamides with bases: 1-amino-6,7,8,9-tetrahydrofuro[2,3-c]isoquinoline-2-carboxamides and 3-amino-4-cyano-5,6,7,8-tetrahydroisoquinolines via a Smiles-type rearrangement. Tetrahedron, 2015, 71, 3263-3272.	1.0	11
82	4,5-Diaryl 3(2H)Furanones: Anti-Inflammatory Activity and Influence on Cancer Growth. Molecules, 2019, 24, 1751.	1.7	11
83	Exploration of the Antimicrobial Effects of Benzothiazolylthiazolidin-4-One and In Silico Mechanistic Investigation. Molecules, 2021, 26, 4061.	1.7	11
84	New Sulfanilamide Derivatives Incorporating Heterocyclic Carboxamide Moieties as Carbonic Anhydrase Inhibitors. Pharmaceuticals, 2021, 14, 828.	1.7	11
85	Study of local anesthetic activity of some derivatives of 3-amino-BENZO-[d]-Isothiazole. SAR and QSAR in Environmental Research, 2003, 14, 485-495.	1.0	10
86	Thiazole Derivatives as Inhibitors of Purified Bovine Liver Mitochondrial Monoamine Oxidase-B: Structure-Activity Relationships and Theoretical Study. Journal of Enzyme Inhibition and Medicinal Chemistry, 1999, 14, 307-321.	0.5	9
87	Synthesis and biological evaluation of lipid-like 5-(2-hydroxyethyl)-4-methyl-1,3-thiazole derivatives as potential anticancer and antimicrobial agents. MedChemComm, 2015, 6, 1464-1470.	3.5	9
88	Inhibition of Renin-Angiotensin System and Advanced Glycation End Products Formation: A Promising Therapeutic Approach Targeting on Cardiovascular Diseases. Cardiovascular and Hematological Agents in Medicinal Chemistry, 2007, 5, 249-264.	0.4	9
89	Carbonic Anhydrase Inhibition with Sulfonamides Incorporating Pyrazole- and Pyridazinecarboxamide Moieties Provides Examples of Isoform-Selective Inhibitors. Molecules, 2021, 26, 7023.	1.7	9
90	Enzymatic Synthesis and Antimicrobial Activity of Oligomer Analogues of Medicinal Biopolymers from Comfrey and Other Species of the Boraginaceae Family. Pharmaceutics, 2022, 14, 115.	2.0	9

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91	Pyrazolo[4,3-c]pyridine Sulfonamides as Carbonic Anhydrase Inhibitors: Synthesis, Biological and In Silico Studies. Pharmaceuticals, 2022, 15, 316.	1.7	9
92	New <i>N</i> -(2-phenyl-4-oxo-1,3-thiazolidin-3-yl)-1,2-benzothiazole-3-carboxamides and acetamides as antimicrobial agents. MedChemComm, 2017, 8, 2142-2154.	3.5	8
93	Derivatives of a new heterocyclic system – pyrano[3,4- <i>c</i>][1,2,4]triazolo[4,3- <i>a</i>]pyridines: synthesis, docking analysis and neurotropic activity. MedChemComm, 2019, 10, 1399-1411.	3.5	8
94	Breakthroughs in Medicinal Chemistry: New Targets and Mechanisms, New Drugs, New Hopes–6. Molecules, 2020, 25, 119.	1.7	8
95	Synthesis and Neurotropic Activity of New Heterocyclic Systems: Pyridofuro[3,2-d]pyrrolo[1,2-a]pyrimidines, Pyridofuro[3,2-d]pyrido[1,2-a]pyrimidines and Pyridofuro[3â€2,2â€2:4,5]pyrimido[1,2-a]azepines. Molecules, 2021, 26, 3320.	1.7	8
96	Discovery of benzothiazole-based thiazolidinones as potential anti-inflammatory agents: anti-inflammatory activity, soybean lipoxygenase inhibition effect and molecular docking studies. SAR and QSAR in Environmental Research, 2022, 33, 485-497.	1.0	8
97	New Methods for the Synthesis of 3â€Aminoâ€6,7â€Dihydroâ€5 <i>H</i> â€Cyclopenta[<i>c</i>]Pyridineâ€4â€Carbonitriles and Cyclopenta[<i>d</i>]Pyrazolo[3,4â€ <i>b</i>]Pyridines via a Smilesâ€type Rearrangement. Journal of Heterocyclic Chemistry, 2017, 54, 1199-1209	1.4	7
98	Synthesis of New Heterocyclic Systems: Pyrido[3′,2′:4,5]thieno(furo)[2,3â€ <i>e</i>][1,2,4]triazolopyrimidines and an Unusual ANRORC Rearrangement in the Fused Pyrimidine Series. ChemistrySelect, 2018, 3, 10938-10942.	0.7	7
99	Synthesis and antimicrobial activity of new 2â€piperazinâ€lâ€ylâ€ <i>N</i> â€l,3â€thiazolâ€2â€ylacetamides of cyclopenta[<i>c</i>]pyridines and pyrano[3,4â€ <i>c</i>]pyridines. Archiv Der Pharmazie, 2021, 354, e2000208.	2.1	7
100	Chromenol Derivatives as Novel Antifungal Agents: Synthesis, In Silico and In Vitro Evaluation. Molecules, 2021, 26, 4304.	1.7	7
101	Synthesis and Evaluation of Antimicrobial Activity and Molecular Docking of New N-1,3-thiazol-2-ylacetamides of Condensed Pyrido[3',2':4,5] furo(thieno)[3,2-d]pyrimidines. Current Topics in Medicinal Chemistry, 2020, 20, 2192-2209.	1.0	7
102	Thiazolyl and Isothiazolyl Azomethine Derivatives with Anti-inflammatory and Antioxidant Activities. Arzneimittelforschung, 2004, 54, 530-537.	0.5	6
103	Synthesis and anti-inflammatory activity of ethynylthiazoles. Chemistry of Heterocyclic Compounds, 2006, 42, 675-680.	0.6	6
104	On the reactivity of 4-cyano-1,3-dichloro-7-methyl-5,6,7,8-tetrahydro-2,7-naphthyridine with several amines in different experimental conditions: monosubstitution, disubstitution, and a new unexpected rearrangement. Tetrahedron, 2014, 70, 4891-4902.	1.0	6
105	Pyridofuropyrrolo[1,2-a]pyrimidines and pyridofuropyrimido[1,2-a]azepines: new chemical entities (NCE) with anticonvulsive and psychotropic properties. RSC Advances, 2016, 6, 32234-32244.	1.7	6
106	2, 2′â€Dihydroxybenzophenones and Derivatives. Efficient Synthesis and Structure Endoscopy by DFT and NMR. Credentials as Potent Antiinflammatory Agents ChemistrySelect, 2016, 1, 2426-2438.	0.7	6
107	Cytotoxicity and Anti–inflammation Profiles of Synthesized Thiazolesâ€Based <i>N</i> â€Bisphosphonates and Relevant Bisphosphonic acids. ChemistrySelect, 2016, 1, 3797-3803.	0.7	6
108	Synthesis, characterization and biological evaluation of Pd(ii), Cu(ii), Re(i) and 99mTc(i) thiazole-based complexes. MedChemComm, 2018, 9, 831-842.	3.5	6

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109	Extending the Inhibition Profiles of Coumarin-Based Compounds Against Human Carbonic Anhydrases: Synthesis, Biological, and In Silico Evaluation. Molecules, 2019, 24, 3580.	1.7	6
110	Chromene-Containing Aromatic Sulfonamides with Carbonic Anhydrase Inhibitory Properties. International Journal of Molecular Sciences, 2021, 22, 5082.	1.8	6
111	Synthesis of 3,3-dimethyl-6-oxopyrano[3,4- <i>c</i>]pyridines and their antiplatelet and vasodilatory activity. Journal of Pharmacy and Pharmacology, 2022, 74, 887-895.	1.2	6
112	Synthesis, In Silico and In Vitro Evaluation. Pharmaceuticals, 2021, 14, 1096.	1.7	6
113	Synthesis, Biological Evaluation and Molecular Docking Studies of 5-IndolyImethylen-4-oxo-2-thioxothiazolidine Derivatives. Molecules, 2022, 27, 1068.	1.7	6
114	Synthesis and structure of a new heterocyclic system: pyrido[3′,2′:4,5]furo[3,2-d][1,2,4]triazolo[4,3-a]pyrimidin-7(8)-one. Tetrahedron Letters, 2016, 57, 5338-5	3407	5
115	Breakthroughs in Medicinal Chemistry: New Targets and Mechanisms, New Drugs, New Hopes–7. Molecules, 2020, 25, 2968.	1.7	5
116	Synthesis of New Derivatives of Heterocyclic Systems Containing Triazolopyrimidine, thiazolo[3,2-a]pyrimidine and pyrimido[2,1-b] thiazine Moiety Showing Promising Antimicrobial Activity. Current Organic Chemistry, 2019, 22, 2576-2588.	0.9	5
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