

Ali Mm

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/4756365/ali-mm-publications-by-year.pdf>

Version: 2024-04-25

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

68
papers

1,597
citations

23
h-index

37
g-index

69
ext. papers

1,890
ext. citations

3.8
avg, IF

4.9
L-index

#	Paper	IF	Citations
68	Anticancer Effects with Molecular Docking Confirmation of Newly Synthesized Isatin Sulfonamide Molecular Hybrid Derivatives against Hepatic Cancer Cell Lines.. <i>Biomedicines</i> , 2022 , 10,	4.8	3
67	Non-ulcerogenic pyrazolyl 2-hydroxychalcones and pyrazolylpyrazolines derived from naturally existing furochromone (khellin): semi-synthesis, docking study and anti-inflammatory activity. <i>Natural Product Research</i> , 2021 , 1-8	2.3	
66	Effective Pharmacophore for CDC25 Phosphatases Enzyme Inhibitors: Newly Synthesized Bromothiazolopyrimidine Derivatives. <i>Mini-Reviews in Medicinal Chemistry</i> , 2021 , 21, 118-131	3.2	3
65	Targeting Receptor Tyrosine Kinase VEGFR-2 in Hepatocellular Cancer: Rational Design, Synthesis and Biological Evaluation of 1,2-Disubstituted Benzimidazoles. <i>Molecules</i> , 2020 , 25,	4.8	13
64	Synthesis and molecular docking study of new pyrazole derivatives as potent anti-breast cancer agents targeting VEGFR-2 kinase. <i>Bioorganic Chemistry</i> , 2020 , 101, 103916	5.1	18
63	Design, synthesis, and molecular docking of novel 2-arylbenzothiazole multiangiokinase inhibitors targeting breast cancer. <i>Archiv Der Pharmazie</i> , 2020 , 353, e1900340	4.3	11
62	Novel potent substituted 4-amino-2-thiopyrimidines as dual VEGFR-2 and BRAF kinase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019 , 179, 707-722	6.8	21
61	Synthesis and In Vitro Antitumor Activity of Novel Chromenones Bearing Benzothiazole Moiety. <i>Russian Journal of Bioorganic Chemistry</i> , 2019 , 45, 42-53	1	16
60	Characterization of a new efficient low molecular weight Bacillus subtilis NRC levansucrase and its levan. <i>Journal of Basic Microbiology</i> , 2019 , 59, 1004-1015	2.7	12
59	Towards breast cancer targeting: Synthesis of tetrahydroindolocarbazoles, antibreast cancer evaluation, uPA inhibition, molecular genetic and molecular modelling studies. <i>Bioorganic Chemistry</i> , 2019 , 93, 103332	5.1	1
58	New 2,4-disubstituted-2-thiopyrimidines as VEGFR-2 inhibitors: Design, synthesis, and biological evaluation. <i>Archiv Der Pharmazie</i> , 2019 , 352, e1900089	4.3	7
57	ZnO Nanoparticles Catalyst in the Synthesis of Bioactive Fused Pyrimidines as Anti-breast Cancer Agents Targeting VEGFR-2. <i>Medicinal Chemistry</i> , 2019 , 15, 277-286	1.8	22
56	Type IIA - Type IIB protein tyrosine kinase inhibitors hybridization as an efficient approach for potent multikinase inhibitor development: Design, synthesis, anti-proliferative activity, multikinase inhibitory activity and molecular modeling of novel indolinone-based ureides and amides. <i>European Journal of Medicinal Chemistry</i> , 2019 , 178, 27-53	6.8	37
55	Synthesis, anticancer effect and molecular modeling of new thiazolylpyrazolyl coumarin derivatives targeting VEGFR-2 kinase and inducing cell cycle arrest and apoptosis. <i>Bioorganic Chemistry</i> , 2019 , 85, 253-273	5.1	38
54	Design, synthesis, and molecular docking of novel indole scaffold-based VEGFR-2 inhibitors as targeted anticancer agents. <i>Archiv Der Pharmazie</i> , 2018 , 351, 1700299	4.3	11
53	The prophylactic and therapeutic effects of Momordica charantia methanol extract through controlling different hallmarks of the hepatocarcinogenesis. <i>Biomedicine and Pharmacotherapy</i> , 2018 , 98, 491-498	7.5	10
52	Anti-proliferative activity of newly synthesized Cd(II), Cu(II), Zn(II), Ni(II), Co(II), VO(II), and Mn(II) complexes of 2-((4,9-dimethoxy-5-oxo-5H-furo[3,2-g]chromen-6-yl)methylene)hydrazinecarbothioamide on three human cancer cells. <i>Applied Organometallic Chemistry</i> , 2018 , 32, e3936	3.1	6

51	Synthesis and anticancer activity of some novel diethyl {(chromonyl/pyrazolyl) [(4-oxo-2-phenyl-quinazolin-3(4H)-yl)amino]methyl}phosphonates. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2018 , 193, 668-674	1	9
50	Tetrahydroindolocarbazoles (THICZs) as new class of urokinase (uPA) inhibitors: Synthesis, anticancer evaluation, DNA-damage determination, and molecular modelling study. <i>Bioorganic Chemistry</i> , 2018 , 80, 545-554	5.1	4
49	l-Amino acid oxidase from <i>Cerastes vipera</i> snake venom: Isolation, characterization and biological effects on bacteria and tumor cell lines. <i>Toxicon</i> , 2018 , 150, 270-279	2.8	16
48	Novel Nitro-Heterocycles Sugar and Indoles Candidates as Lead Structures Targeting HepG2 and A549 Cancer Cell Lines. <i>Current Bioactive Compounds</i> , 2018 , 14, 434-444	0.9	3
47	Design, Synthesis, In Vitro Anti-cancer Activity, ADMET Profile and Molecular Docking of Novel Triazolo[3,4-a]phthalazine Derivatives Targeting VEGFR-2 Enzyme. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2018 , 18, 1184-1196	2.2	28
46	Antioxidant and anticancer efficacy of therapeutic bioactive compounds from fermented olive waste. <i>Grasas Y Aceites</i> , 2018 , 69, 266	1.3	4
45	Part II: New candidates of pyrazole-benzimidazole conjugates as checkpoint kinase 2 (Chk2) inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2018 , 144, 859-873	6.8	14
44	Benzimidazole - Schiff bases and their complexes: synthesis, anticancer activity and molecular modeling as Aurora kinase inhibitor. <i>Zeitschrift Fur Naturforschung - Section C Journal of Biosciences</i> , 2018 , 73, 465-478	1.7	15
43	Part I: Design, synthesis and biological evaluation of novel pyrazole-benzimidazole conjugates as checkpoint kinase 2 (Chk2) inhibitors with studying their activities alone and in combination with genotoxic drugs. <i>European Journal of Medicinal Chemistry</i> , 2017 , 134, 392-405	6.8	23
42	Design, synthesis, molecular docking and cytotoxic evaluation of novel 2-furybenzimidazoles as VEGFR-2 inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2017 , 136, 315-329	6.8	50
41	Reaction of 2-cyano[(4-oxo-4H-chromen-3-yl)methylidene]acetohydrazide with phosphorus reagents: Synthesis and evaluation of anticancer activities of some novel 1,2-azaphospholes, 1,2,3-diazaphospholidine, and 1,3,2-diaza-phosphinanes bearing a chromone ring. <i>Synthetic Communications</i> , 2017 , 47, 1458-1470	1.7	10
40	Synthesis and Structure-Activity Relationship Study of Novel Pyrazolythiazoles as Potential Anti-Breast Cancer Agents. <i>Journal of Heterocyclic Chemistry</i> , 2017 , 54, 1974-1982	1.9	4
39	New Coumarin Derivatives as Anti-Breast and Anti-Cervical Cancer Agents Targeting VEGFR-2 and p38MAPK. <i>Archiv Der Pharmazie</i> , 2017 , 350, 1700064	4.3	27
38	Design, Synthesis, Molecular Docking, and Anticancer Activity of Phthalazine Derivatives as VEGFR-2 Inhibitors. <i>Archiv Der Pharmazie</i> , 2017 , 350, 1700240	4.3	42
37	Synthesis and cytotoxic activity of certain benzothiazole derivatives against human MCF-7 cancer cell line. <i>Chemical Biology and Drug Design</i> , 2017 , 89, 566-576	2.9	13
36	Synthesis, characterization, and in vitro anticancer evaluation of iron oxide/chitosan nanocomposites. <i>Inorganic and Nano-Metal Chemistry</i> , 2017 , 47, 405-411	1.2	8
35	Synthesis of some novel 4-benzothiazol-2-yl-benzoyl-1H-pyrazoles, and evaluation as antiangiogenic agents. <i>Research on Chemical Intermediates</i> , 2016 , 42, 1521-1536	2.8	13
34	Increasing the binding affinity of VEGFR-2 inhibitors by extending their hydrophobic interaction with the active site: Design, synthesis and biological evaluation of 1-substituted-4-(4-methoxybenzyl)phthalazine derivatives. <i>European Journal of Medicinal Chemistry</i> , 2016 , 113, 50-62	6.8	59

33	Synthesis, characterization, and antitumor evaluation of 4-aminoximidofurazan derivatives. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2016 , 191, 1000-1008	1	8
32	1-Piperazinyolphthalazines as potential VEGFR-2 inhibitors and anticancer agents: Synthesis and in vitro biological evaluation. <i>European Journal of Medicinal Chemistry</i> , 2016 , 107, 165-79	6.8	70
31	Synthesis and Antiproliferative Activities of Benzimidazole-Based Sulfide and Sulfoxide Derivatives. <i>Scientia Pharmaceutica</i> , 2016 , 84, 1-18	4.3	2
30	Synthesis, in vitro and in vivo antitumor and antiviral activity of novel 1-substituted benzimidazole derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 826-45	5.6	28
29	Synthesis and Characterization of New 3,5-DIARYL-3H-Dispiropyran/Thiopyran[4,2'-Chroman-3,2'-[1,3,4-Thiadiazol]-4'-One Derivatives and Related Compounds as Anticancer and Antiviral Agents. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2015 , 190, 1901-1911	1	3
28	Synthesis and antiproliferative activity of novel polynuclear heterocyclic compounds derived from 2,3-diaminophenazine. <i>European Journal of Medicinal Chemistry</i> , 2015 , 90, 568-76	6.8	10
27	Synthesis and in vitro cytotoxic activity of novel pyrazolo[1,5-a]pyrimidines and related Schiff bases. <i>Turkish Journal of Chemistry</i> , 2015 , 39, 1102-1113	1	33
26	Indoline ureas as potential anti-hepatocellular carcinoma agents targeting VEGFR-2: Synthesis, in vitro biological evaluation and molecular docking. <i>European Journal of Medicinal Chemistry</i> , 2015 , 100, 89-97	6.8	46
25	Celecoxib analogs bearing benzofuran moiety as cyclooxygenase-2 inhibitors: design, synthesis and evaluation as potential anti-inflammatory agents. <i>European Journal of Medicinal Chemistry</i> , 2014 , 76, 482-93	6.8	98
24	Synthesis, biological evaluation, and docking studies of new 2-furylbenzimidazoles as anti-angiogenic agents: part II. <i>Archiv Der Pharmazie</i> , 2014 , 347, 291-304	4.3	13
23	Part I. Synthesis, biological evaluation and docking studies of new 2-furylbenzimidazoles as antiangiogenic agents. <i>European Journal of Medicinal Chemistry</i> , 2014 , 87, 868-80	6.8	22
22	Some pyrazole and pyrazolo[3,4-d]pyrimidine derivatives: synthesis and anticancer evaluation. <i>Archiv Der Pharmazie</i> , 2014 , 347, 559-65	4.3	14
21	Design, synthesis and molecular docking study of novel quinoxalin-2(1H)-ones as anti-tumor active agents with inhibition of tyrosine kinase receptor and studying their cyclooxygenase-2 activity. <i>European Journal of Medicinal Chemistry</i> , 2014 , 86, 122-32	6.8	78
20	Synthesis, Characterization, and Antiproliferative Activity of Cu ²⁺ , V(IV)O ₂ ⁺ , Co ²⁺ , Mn ²⁺ , and Ni ²⁺ Complexes with 3-(2-(4-Methoxyphenylcarbamoithiyl)Hydrazinyl)-3-OXO-N-(Thiazol-2-yl)Propanamide against Human Hepatocellular Carcinoma Cells. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2014 , 189, 1001-1011	1	1
19	Photodynamic therapy mediated antiproliferative activity of some metal-doped ZnO nanoparticles in human liver adenocarcinoma HepG2 cells under UV irradiation. <i>Journal of Photochemistry and Photobiology B: Biology</i> , 2014 , 138, 99-108	6.7	35
18	Anticancer evaluation of some newly synthesized N-nicotinonitrile derivative. <i>European Journal of Medicinal Chemistry</i> , 2013 , 69, 521-6	6.8	16
17	Design, synthesis and structure-activity relationship of novel quinoxaline derivatives as cancer chemopreventive agent by inhibition of tyrosine kinase receptor. <i>European Journal of Medicinal Chemistry</i> , 2013 , 69, 115-24	6.8	32
16	Levansucrase optimization using solid state fermentation and levan biological activities studies. <i>Carbohydrate Polymers</i> , 2013 , 96, 332-41	10.3	21

15	Synthesis and docking studies of novel antitumor benzimidazoles. <i>Bioorganic and Medicinal Chemistry</i> , 2012 , 20, 6989-7001	3-4	31
14	Synthesis, characterization and anticancer studies of ferrocenyl complexes containing thiazole moiety. <i>Applied Organometallic Chemistry</i> , 2012 , 26, 230-236	3-1	17
13	New pyrimidinone and fused pyrimidinone derivatives as potential anticancer chemotherapeutics. <i>Archiv Der Pharmazie</i> , 2012 , 345, 729-38	4-3	7
12	Phytochemical investigation and biological studies of <i>Bombax malabaricum</i> flowers. <i>Natural Product Research</i> , 2011 , 25, 141-51	2-3	22
11	Synthesis and in vitro cytotoxic activity of novel pyrazolo[3,4-d]pyrimidines and related pyrazole hydrazones toward breast adenocarcinoma MCF-7 cell line. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 6808-17	3-4	62
10	Synthesis and anticancer effects of some novel pyrazolo[3,4-d]pyrimidine derivatives by generating reactive oxygen species in human breast adenocarcinoma cells. <i>European Journal of Medicinal Chemistry</i> , 2011 , 46, 1019-26	6-8	64
9	Synthesis, Spectral, Characterization, and Anticancer Activity of Some Binary and Mixed Ligand Complexes of 4-Methyl-2-Pentanone Thiosemicarbazone and Some Amino Acids. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2010 , 185, 2171-2181	1	9
8	Synthesis, characterization, and anticancer properties of ferrocenyl complexes containing a salicylaldehyde moiety. <i>Monatshefte Für Chemie</i> , 2010 , 141, 387-393	1-4	17
7	Synthesis of new quinoline derivatives as inhibitors of human tumor cells growth. <i>Archiv Der Pharmazie</i> , 2010 , 343, 440-8	4-3	35
6	Modulation of anticancer drug-induced P-glycoprotein expression by naringin. <i>Zeitschrift Fur Naturforschung - Section C Journal of Biosciences</i> , 2009 , 64, 109-16	1-7	13
5	Synthesis and in vitro antitumor activity of new substituted thiopyrimidine acyclic nucleosides and their thioglycoside analogs. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2009 , 28, 261-74	1-4	36
4	Amelioration of streptozotocin-induced diabetes mellitus, oxidative stress and dyslipidemia in rats by tomato extract lycopene. <i>Scandinavian Journal of Clinical and Laboratory Investigation</i> , 2009 , 69, 371-9 ²		72
3	Synthesis and Antitumor Evaluation of Some Newly Synthesized Pyrazolopyrimidine and Pyrazolotriazolopyrimidine Derivatives. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2009 , 185, 74-83	1	6
2	The influence of naringin on the oxidative state of rats with streptozotocin-induced acute hyperglycaemia. <i>Zeitschrift Fur Naturforschung - Section C Journal of Biosciences</i> , 2004 , 59, 726-33	1-7	48
1	Induction of metallothionein by zinc protects from daunorubicin toxicity in rats. <i>Toxicology</i> , 2002 , 179, 85-93	4-4	56