Giovanni Marzaro

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46
papers

S10
citations

h-index

27
g-index

52
ext. papers

5 4.12
ext. papers

avg, IF

L-index

#	Paper	IF	Citations
46	Quinazoline derivatives as potential anticancer agents: a patent review (2007 - 2010). <i>Expert Opinion on Therapeutic Patents</i> , 2012 , 22, 223-52	6.8	86
45	Using the TOPS-MODE approach to fit multi-target QSAR models for tyrosine kinases inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2011 , 46, 2185-92	6.8	51
44	Synthesis and antitumor activity of novel amsacrine analogs: the critical role of the acridine moiety in determining their biological activity. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 523-9	3.4	50
43	Quinazoline-based multi-tyrosine kinase inhibitors: synthesis, modeling, antitumor and antiangiogenic properties. <i>European Journal of Medicinal Chemistry</i> , 2013 , 67, 373-83	6.8	49
42	Exploring epidermal growth factor receptor (EGFR) inhibitor features: the role of fused dioxygenated rings on the quinazoline scaffold. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 1862-6	8.3	46
41	Third-generation CDK inhibitors: A review on the synthesis and binding modes of Palbociclib, Ribociclib and Abemaciclib. <i>European Journal of Medicinal Chemistry</i> , 2019 , 172, 143-153	6.8	44
40	Microwave-promoted mono-N-alkylation of aromatic amines in water: a new efficient and green method for an old and problematic reaction. <i>Green Chemistry</i> , 2009 , 11, 774	10	35
39	Development of a novel furocoumarin derivative inhibiting NF- B dependent biological functions: design, synthesis and biological effects. <i>European Journal of Medicinal Chemistry</i> , 2011 , 46, 4870-7	6.8	32
38	A novel approach to quinazolin-4(3H)-one via quinazoline oxidation: an improved synthesis of 4-anilinoquinazolines. <i>Tetrahedron</i> , 2010 , 66, 962-968	2.4	29
37	Psoralen derivatives as inhibitors of NF- B /DNA interaction: synthesis, molecular modeling, 3D-QSAR, and biological evaluation. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 1830-42	8.3	28
36	Lejeuneaceae (Marchantiophyta) from a species-rich taphocoenosis in Miocene Mexican amber, with a review of liverworts fossilised in amber. <i>Review of Palaeobotany and Palynology</i> , 2015 , 221, 59-70) ^{1.7}	26
35	A microwave improvement in the synthesis of the quinazoline scaffold. <i>Tetrahedron Letters</i> , 2007 , 48, 3229-3231	2	26
34	Molecular Mechanism of Action of Trimethylangelicin Derivatives as CFTR Modulators. <i>Frontiers in Pharmacology</i> , 2018 , 9, 719	5.6	22
33	Discovery of biarylaminoquinazolines as novel tubulin polymerization inhibitors. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 4598-4605	8.3	21
32	Targeting kinases with anilinopyrimidines: discovery of N-phenyl-NS[4-(pyrimidin-4-ylamino)phenyl]urea derivatives as selective inhibitors of class III receptor tyrosine kinase subfamily. <i>Scientific Reports</i> , 2015 , 5, 16750	4.9	21
31	A novel convenient synthesis of benzoquinazolines. <i>Organic Letters</i> , 2006 , 8, 255-6	6.2	20
30	Known Triterpenes and their Derivatives as Scaffolds for the Development of New Therapeutic Agents for Cancer. <i>Current Medicinal Chemistry</i> , 2018 , 25, 1259-1269	4.3	20

(2015-2016)

29	Investigational drugs targeting cyclin-dependent kinases for the treatment of cancer: an update on recent findings (2013-2016). <i>Expert Opinion on Investigational Drugs</i> , 2016 , 25, 1215-30	5.9	19
28	A new access to quinazolines from simple anilines. <i>Tetrahedron</i> , 2006 , 62, 12351-12356	2.4	19
27	Benzoquinazoline derivatives as new agents affecting DNA processing. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 1197-204	3.4	17
26	The Moss Macromitrium Richardii (Orthotrichaceae) with Sporophyte and Calyptra Enclosed in Hymenaea Resin from the Dominican Republic. <i>Polish Botanical Journal</i> , 2013 , 58, 221-230		14
25	Structural and Functional Insights on an Uncharacterized AEGlobin-Gene Polymorphism Present in Four D-Thalassemia Families with High Fetal Hemoglobin Levels. <i>Molecular Diagnosis and Therapy</i> , 2016 , 20, 161-73	4.5	13
24	Differential Effects of Angelicin Analogues on NF-B Activity and IL-8 Gene Expression in Cystic Fibrosis IB3-1 Cells. <i>Mediators of Inflammation</i> , 2017 , 2017, 2389487	4.3	12
23	Corilagin Induces High Levels of Apoptosis in the Temozolomide-Resistant T98G Glioma Cell Line. <i>Oncology Research</i> , 2018 , 26, 1307-1315	4.8	12
22	Substituted quinazolinones as kinase inhibitors endowed with anti-fibrotic properties. <i>European Journal of Medicinal Chemistry</i> , 2016 , 115, 416-25	6.8	11
21	Pyrroloquinolinone-based dual topoisomerase I/II inhibitor. <i>European Journal of Medicinal Chemistry</i> , 2014 , 77, 103-9	6.8	9
20	Early Evaluation of Copper Radioisotope Production at ISOLPHARM. <i>Molecules</i> , 2018 , 23,	4.8	9
19	Photochemical and photobiological studies on furoquinazolines as new psoralen analogs. <i>Journal of Photochemistry and Photobiology B: Biology</i> , 2014 , 138, 43-54	6.7	7
19		6. ₇	7
	Photochemistry and Photobiology B: Biology, 2014 , 138, 43-54 Targeting DNA Binding for NF- B as an Anticancer Approach in Hepatocellular Carcinoma. <i>Cells</i> ,		7 7 6
18	Photochemistry and Photobiology B: Biology, 2014, 138, 43-54 Targeting DNA Binding for NF-B as an Anticancer Approach in Hepatocellular Carcinoma. <i>Cells</i> , 2018, 7, Design, synthesis and biological evaluation of novel trimethylangelicin analogues targeting nuclear	7.9	7
18	Photochemistry and Photobiology B: Biology, 2014, 138, 43-54 Targeting DNA Binding for NF-B as an Anticancer Approach in Hepatocellular Carcinoma. Cells, 2018, 7, Design, synthesis and biological evaluation of novel trimethylangelicin analogues targeting nuclear factor kB (NF-kB). European Journal of Medicinal Chemistry, 2018, 151, 285-293 Novel benzoquinoline derivatives via unpredicted condensation of ethyl propiolate and naphthylamines: Synthesis and topoisomerase inhibition activity. Bioorganic and Medicinal	7.9 6.8	7
18 17 16	Photochemistry and Photobiology B: Biology, 2014, 138, 43-54 Targeting DNA Binding for NF-B as an Anticancer Approach in Hepatocellular Carcinoma. Cells, 2018, 7, Design, synthesis and biological evaluation of novel trimethylangelicin analogues targeting nuclear factor kB (NF-kB). European Journal of Medicinal Chemistry, 2018, 151, 285-293 Novel benzoquinoline derivatives via unpredicted condensation of ethyl propiolate and naphthylamines: Synthesis and topoisomerase inhibition activity. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 4875-4878 New vandetanib analogs: fused tricyclic quinazolines with antiangiogenic potential. Investigational	7.9 6.8 2.9	7 6
18 17 16 15	Photochemistry and Photobiology B: Biology, 2014, 138, 43-54 Targeting DNA Binding for NF-B as an Anticancer Approach in Hepatocellular Carcinoma. Cells, 2018, 7, Design, synthesis and biological evaluation of novel trimethylangelicin analogues targeting nuclear factor kB (NF-kB). European Journal of Medicinal Chemistry, 2018, 151, 285-293 Novel benzoquinoline derivatives via unpredicted condensation of ethyl propiolate and naphthylamines: Synthesis and topoisomerase inhibition activity. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 4875-4878 New vandetanib analogs: fused tricyclic quinazolines with antiangiogenic potential. Investigational New Drugs, 2012, 30, 594-603	7.9 6.8 2.9	7 6 6

11	QSAR and 3D-QSAR models in the field of tubulin inhibitors as anticancer agents. <i>Current Topics in Medicinal Chemistry</i> , 2014 , 14, 2253-62	3	4
10	Toward novel sulphur-containing derivatives of tetraazacyclododecane: synthesis, acidBase properties, spectroscopic characterization, DFT calculations, and cadmium(II) complex formation in aqueous solution. <i>New Journal of Chemistry</i> , 2020 , 44, 8337-8350	3.6	4
9	Photobiological properties of 3-psoralenacetic acids. <i>Photochemical and Photobiological Sciences</i> , 2015 , 14, 2074-86	4.2	2
8	Autogrid-based clustering of kinases: selection of representative conformations for docking purposes. <i>Molecular Diversity</i> , 2014 , 18, 611-9	3.1	2
7	Psoralenquinones as a novel class of proteasome inhibitors: design, synthesis and biological evaluation. <i>ChemMedChem</i> , 2011 , 6, 996-1000	3.7	2
6	Preliminary evaluation of the production of non-carrier added Ag as core of a therapeutic radiopharmaceutical in the framework of ISOLPHARM_Ag experiment. <i>Applied Radiation and Isotopes</i> , 2020 , 164, 109258	1.7	2
5	Preliminary Study of a 1,5-Benzodiazepine-Derivative Labelled with Indium-111 for CCK-2 Receptor Targeting. <i>Molecules</i> , 2021 , 26,	4.8	2
4	The Importance of Descriptor-Based Clusterization in QSAR Models Development: Tyrosine Kinases Inhibitors as a Key Study. <i>Molecular Informatics</i> , 2011 , 30, 721-32	3.8	1
3	Tyrosine kinase inhibitor prodrug-loaded liposomes for controlled release at tumor microenvironment. <i>Journal of Controlled Release</i> , 2021 , 340, 318-330	11.7	1
2	Dual Kinase Targeting in Leukemia. <i>Cancers</i> , 2021 , 13,	6.6	1
1	Surface plasmon resonance based analysis of the binding of LYAR protein to the rs368698783 (G>A) polymorphic Alglobin gene sequences mutated in Ethalassemia. <i>Analytical and Bioanalytical Chemistry</i> , 2019 , 411, 7699-7707	4.4	O