

Giovanni Marzaro

List of Publications by Year in descending order

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47
papers

1,076
citations

448610

19
h-index

466096

32
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52
all docs

52
docs citations

52
times ranked

1954
citing authors

#	ARTICLE	IF	CITATIONS
1	Proteolysis Targeting Chimeric Molecules: Tuning Molecular Strategies for a Clinically Sound Listening. <i>International Journal of Molecular Sciences</i> , 2022, 23, 6630.	1.8	8
2	Dual Kinase Targeting in Leukemia. <i>Cancers</i> , 2021, 13, 119.	1.7	2
3	Preliminary Study of a 1,5-Benzodiazepine-Derivative Labelled with Indium-111 for CCK-2 Receptor Targeting. <i>Molecules</i> , 2021, 26, 918.	1.7	8
4	Tyrosine kinase inhibitor prodrug-loaded liposomes for controlled release at tumor microenvironment. <i>Journal of Controlled Release</i> , 2021, 340, 318-330.	4.8	8
5	Preliminary evaluation of the production of non-carrier added ¹¹¹ Ag as core of a therapeutic radiopharmaceutical in the framework of ISOLPHARM_Ag experiment. <i>Applied Radiation and Isotopes</i> , 2020, 164, 109258.	0.7	10
6	Toward novel sulphur-containing derivatives of tetraazacyclododecane: synthesis, acid-base properties, spectroscopic characterization, DFT calculations, and cadmium(II) complex formation in aqueous solution. <i>New Journal of Chemistry</i> , 2020, 44, 8337-8350.	1.4	11
7	Surface plasmon resonance based analysis of the binding of LYAR protein to the rs368698783 (G>A) polymorphic A β -globin gene sequences mutated in β ² -thalassemia. <i>Analytical and Bioanalytical Chemistry</i> , 2019, 411, 7699-7707.	1.9	1
8	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.	2.6	72
9	Design, synthesis and biological evaluation of novel trimethylangelicin analogues targeting nuclear factor κ B (NF- κ B). <i>European Journal of Medicinal Chemistry</i> , 2018, 151, 285-293.	2.6	13
10	Corilagin Induces High Levels of Apoptosis in the Temozolomide-Resistant T98G Glioma Cell Line. <i>Oncology Research</i> , 2018, 26, 1307-1315.	0.6	18
11	Early Evaluation of Copper Radioisotope Production at ISOLPHARM. <i>Molecules</i> , 2018, 23, 2437.	1.7	16
12	Targeting DNA Binding for NF- κ B as an Anticancer Approach in Hepatocellular Carcinoma. <i>Cells</i> , 2018, 7, 177.	1.8	11
13	Molecular Mechanism of Action of Trimethylangelicin Derivatives as CFTR Modulators. <i>Frontiers in Pharmacology</i> , 2018, 9, 719.	1.6	28
14	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.	1.2	30
15	Multicomponent Assembly of the Kinesin Spindle Protein Inhibitor CPLYJ039 and Analogues as Antimitotic Agents. <i>ACS Combinatorial Science</i> , 2017, 19, 153-160.	3.8	5
16	Discovery of ^{wt} RET and ^{V804M} RET Inhibitors: From Hit to Lead. <i>ChemMedChem</i> , 2017, 12, 1390-1398.	1.6	7
17	Differential Effects of Angelicin Analogues on NF- κ B Activity and IL-8 Gene Expression in Cystic Fibrosis IB3-1 Cells. <i>Mediators of Inflammation</i> , 2017, 2017, 1-11.	1.4	16
18	Substituted quinazolinones as kinase inhibitors endowed with anti-fibrotic properties. <i>European Journal of Medicinal Chemistry</i> , 2016, 115, 416-425.	2.6	12

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19	Novel benzoquinoline derivatives via unpredicted condensation of ethyl propiolate and naphthylamines: Synthesis and topoisomerase inhibition activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 4875-4878.	1.0	6
20	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, 1215-1230.	1.9	24
21	Structural and Functional Insights on an Uncharacterized $\text{A}^{\hat{1}3}$ -Globin-Gene Polymorphism Present in Four $\text{I}^{\hat{2}0}$ -Thalassemia Families with High Fetal Hemoglobin Levels. <i>Molecular Diagnosis and Therapy</i> , 2016, 20, 161-173.	1.6	17
22	Targeting kinases with anilinopyrimidines: discovery of N-phenyl-N \hat{e} TM -[4-(pyrimidin-4-ylamino)phenyl]urea derivatives as selective inhibitors of class III receptor tyrosine kinase subfamily. <i>Scientific Reports</i> , 2015, 5, 16750.	1.6	53
23	Psoralen derivatives as inhibitors of NF- $\hat{\kappa}$ B/DNA interaction: the critical role of the furan ring. <i>Molecular Diversity</i> , 2015, 19, 551-561.	2.1	6
24	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.	0.8	36
25	Photobiological properties of 3-psoralenacetic acids. <i>Photochemical and Photobiological Sciences</i> , 2015, 14, 2074-2086.	1.6	3
26	Pyrroloquinolinone-based dual topoisomerase I/II inhibitor. <i>European Journal of Medicinal Chemistry</i> , 2014, 77, 103-109.	2.6	13
27	Autogrid-based clustering of kinases: selection of representative conformations for docking purposes. <i>Molecular Diversity</i> , 2014, 18, 611-619.	2.1	3
28	Discovery of Biaryl aminoquinazolines as Novel Tubulin Polymerization Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 4598-4605.	2.9	28
29	Photochemical and photobiological studies on furoquinazolines as new psoralen analogs. <i>Journal of Photochemistry and Photobiology B: Biology</i> , 2014, 138, 43-54.	1.7	9
30	QSAR and 3D-QSAR Models in the Field of Tubulin Inhibitors as Anticancer Agents. <i>Current Topics in Medicinal Chemistry</i> , 2014, 14, 2253-2262.	1.0	7
31	Quinazoline-based multi-tyrosine kinase inhibitors: Synthesis, modeling, antitumor and antiangiogenic properties. <i>European Journal of Medicinal Chemistry</i> , 2013, 67, 373-383.	2.6	59
32	Psoralen Derivatives as Inhibitors of NF- $\hat{\kappa}$ B/DNA Interaction: Synthesis, Molecular Modeling, 3D-QSAR, and Biological Evaluation. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 1830-1842.	2.9	34
33	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.	0.5	17
34	Quinazoline derivatives as potential anticancer agents: a patent review (2007 \hat{e} 2010). <i>Expert Opinion on Therapeutic Patents</i> , 2012, 22, 223-252.	2.4	104
35	New Vandetanib analogs: fused tricyclic quinazolines with antiangiogenic potential. <i>Investigational New Drugs</i> , 2012, 30, 594-603.	1.2	7
36	Development of a novel furocoumarin derivative inhibiting NF- $\hat{\kappa}$ B dependent biological functions: Design, synthesis and biological effects. <i>European Journal of Medicinal Chemistry</i> , 2011, 46, 4870-4877.	2.6	38

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37	Benzoquinazoline derivatives as new agents affecting DNA processing. <i>Bioorganic and Medicinal Chemistry</i> , 2011, 19, 1197-1204.	1.4	18
38	The Importance of Descriptor-Based Clusterization in QSAR Models Development: Tyrosine Kinases Inhibitors as a Key Study. <i>Molecular Informatics</i> , 2011, 30, 721-732.	1.4	1
39	Psoralenquinones as a Novel Class of Proteasome Inhibitors: Design, Synthesis and Biological Evaluation. <i>ChemMedChem</i> , 2011, 6, 996-1000.	1.6	3
40	Using the TOPS-MODE approach to fit multi-target QSAR models for tyrosine kinases inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2011, 46, 2185-2192.	2.6	62
41	A novel approach to quinazolin-4(3H)-one via quinazoline oxidation: an improved synthesis of 4-anilinoquinazolines. <i>Tetrahedron</i> , 2010, 66, 962-968.	1.0	34
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, 1862-1866.	2.9	51
43	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, 523-529.	1.4	56
44	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.	4.6	39
45	A microwave improvement in the synthesis of the quinazoline scaffold. <i>Tetrahedron Letters</i> , 2007, 48, 3229-3231.	0.7	27
46	A Novel Convenient Synthesis of Benzoquinazolines. <i>Organic Letters</i> , 2006, 8, 255-256.	2.4	22
47	A new access to quinazolines from simple anilines. <i>Tetrahedron</i> , 2006, 62, 12351-12356.	1.0	21