

Elena Mariotto

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4292132/publications.pdf>

Version: 2024-02-01

30
papers

499
citations

623734

14
h-index

713466

21
g-index

31
all docs

31
docs citations

31
times ranked

887
citing authors

#	ARTICLE	IF	CITATIONS
1	Cinnamic acid derivatives linked to arylpiperazines as novel potent inhibitors of tyrosinase activity and melanin synthesis. <i>European Journal of Medicinal Chemistry</i> , 2022, 231, 114147.	5.5	18
2	Biological Evaluation of New Thienopyridinium and Thienopyrimidinium Derivatives as Human Choline Kinase Inhibitors. <i>Pharmaceutics</i> , 2022, 14, 715.	4.5	2
3	Histone Deacetylase Inhibitors Impair Glioblastoma Cell Motility and Proliferation. <i>Cancers</i> , 2022, 14, 1897.	3.7	11
4	Synthesis and Biological Evaluation of Highly Active 7-Anilino Triazolopyrimidines as Potent Antimicrotubule Agents. <i>Pharmaceutics</i> , 2022, 14, 1191.	4.5	7
5	A facile synthesis of diaryl pyrroles led to the discovery of potent colchicine site antimitotic agents. <i>European Journal of Medicinal Chemistry</i> , 2021, 214, 113229.	5.5	13
6	Developing novel classes of protein kinase CK1 \hat{I} inhibitors by fusing [1,2,4]triazole with different bicyclic heteroaromatic systems. <i>European Journal of Medicinal Chemistry</i> , 2021, 216, 113331.	5.5	9
7	Anticancer and Structure Activity Relationship of Non-Symmetrical Choline Kinase Inhibitors. <i>Pharmaceutics</i> , 2021, 13, 1360.	4.5	3
8	BAG1 down \hat{I} regulation increases chemo \hat{I} sensitivity of acute lymphoblastic leukaemia cells. <i>Journal of Cellular and Molecular Medicine</i> , 2021, 25, 9060-9065.	3.6	3
9	Synthesis, biological evaluation, in silico modeling and crystallization of novel small monocationic molecules with potent antiproliferative activity by dual mechanism. <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112797.	5.5	4
10	Autophagic flux inhibition enhances cytotoxicity of the receptor tyrosine kinase inhibitor ponatinib. <i>Journal of Experimental and Clinical Cancer Research</i> , 2020, 39, 195.	8.6	12
11	Design, synthesis, in \hat{I} vitro and in \hat{I} vivo biological evaluation of 2-amino-3-arylbenzo[b]furan derivatives as highly potent tubulin polymerization inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2020, 200, 112448.	5.5	25
12	A BAG's life: Every connection matters in cancer. , 2020, 209, 107498.		26
13	Design, synthesis and biological evaluation of novel vicinal diaryl-substituted 1H-Pyrazole analogues of combretastatin A-4 as highly potent tubulin polymerization inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019, 181, 111577.	5.5	22
14	Design, Synthesis, and Biological Evaluation of 6-Substituted Thieno[3,2- <i>d</i>]pyrimidine Analogues as Dual Epidermal Growth Factor Receptor Kinase and Microtubule Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 1274-1290.	6.4	33
15	Potent In Vitro and In Vivo Anticancer Activity of New Bipyridine and Bipyrimidine Gold (III) Dithiocarbamate Derivatives. <i>Cancers</i> , 2019, 11, 474.	3.7	41
16	SYK Targeting Represents a Potential Therapeutic Option for Relapsed Resistant Pediatric ETV6-RUNX1 B-Acute Lymphoblastic Leukemia Patients. <i>International Journal of Molecular Sciences</i> , 2019, 20, 6175.	4.1	10
17	Abstract A058: Dielectric characterization of glioblastoma cancer stem cells. , 2019, , .		0
18	Abstract A066: Role of the NRF2 signaling pathway in sustaining chemoresistance in medulloblastoma. , 2019, , .		0

#	ARTICLE	IF	CITATIONS
19	TP-0903 inhibits neuroblastoma cell growth and enhances the sensitivity to conventional chemotherapy. <i>European Journal of Pharmacology</i> , 2018, 818, 435-448.	3.5	29
20	AKR1C enzymes sustain therapy resistance in paediatric T-ALL. <i>British Journal of Cancer</i> , 2018, 118, 985-994.	6.4	31
21	Choline Kinase Alpha Inhibition by EB-3D Triggers Cellular Senescence, Reduces Tumor Growth and Metastatic Dissemination in Breast Cancer. <i>Cancers</i> , 2018, 10, 391.	3.7	23
22	Lead optimization-hit expansion of new asymmetrical pyridinium/quinolinium compounds as choline kinase $\hat{1}$ inhibitors. <i>Future Medicinal Chemistry</i> , 2018, 10, 1769-1786.	2.3	4
23	EB-3D a novel choline kinase inhibitor induces deregulation of the AMPK-mTOR pathway and apoptosis in leukemia T-cells. <i>Biochemical Pharmacology</i> , 2018, 155, 213-223.	4.4	19
24	Synthesis and Biological Evaluation of 2-Methyl-4,5-Disubstituted Oxazoles as a Novel Class of Highly Potent Antitubulin Agents. <i>Scientific Reports</i> , 2017, 7, 46356.	3.3	17
25	HLA Class II Antibody Activation of Endothelial Cells Promotes Th17 and Disrupts Regulatory T Lymphocyte Expansion. <i>American Journal of Transplantation</i> , 2016, 16, 1408-1420.	4.7	72
26	Design, synthesis, crystallization and biological evaluation of new symmetrical biscationic compounds as selective inhibitors of human Choline Kinase $\hat{1}$ (Chok $\hat{1}$). <i>Scientific Reports</i> , 2016, 6, 23793.	3.3	21
27	Design and Synthesis of Potent in Vitro and in Vivo Anticancer Agents Based on 1-(3,4,5-Trimethoxyphenyl)-2-Aryl-1H-Imidazole. <i>Scientific Reports</i> , 2016, 6, 26602.	3.3	29
28	Abstract 1297: CDK4/CDK6 inhibition in childhood B-acute lymphoblastic leukemia: a new strategy to mediate glucocorticoid sensitivity. , 2016, , .		1
29	Abstract 1233: In vitro and in vivo pharmacological study of EB-3D: a novel choline kinase inhibitor for breast cancer treatment. , 2016, , .		0
30	New more polar symmetrical bipyridinic compounds: new strategy for the inhibition of choline kinase $\hat{1}$. <i>Future Medicinal Chemistry</i> , 2015, 7, 417-436.	2.3	14