

Mariangela Agamennone

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/94826/publications.pdf>

Version: 2024-02-01

41
papers

757
citations

471061

17
h-index

552369

26
g-index

42
all docs

42
docs citations

42
times ranked

1151
citing authors

#	ARTICLE	IF	CITATIONS
1	Structural Insight into the Stereoselective Inhibition of MMP-8 by Enantiomeric Sulfonamide Phosphonates. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 923-931.	2.9	70
2	Bovine lactoferrin-derived peptides as novel broad-spectrum inhibitors of influenza virus. <i>Pathogens and Global Health</i> , 2012, 106, 12-19.	1.0	53
3	N-Hydroxyurea as zinc binding group in matrix metalloproteinase inhibition: Mode of binding in a complex with MMP-8. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006, 16, 20-24.	1.0	52
4	Biphenyl Sulfonylamino Methyl Bisphosphonic Acids as Inhibitors of Matrix Metalloproteinases and Bone Resorption. <i>ChemMedChem</i> , 2011, 6, 1258-1268.	1.6	44
5	±-Biphenylsulfonylamino 2-methylpropyl phosphonates: Enantioselective synthesis and selective inhibition of MMPs. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 791-799.	1.4	39
6	Arylamino methylene bisphosphonate derivatives as bone seeking matrix metalloproteinase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 6456-6465.	1.4	34
7	Bovine Lactoferrin Prevents Influenza A Virus Infection by Interfering with the Fusogenic Function of Viral Hemagglutinin. <i>Viruses</i> , 2019, 11, 51.	1.5	33
8	AMBER force field implementation of the boronate function to simulate the inhibition of β-lactamases by alkyl and aryl boronic acids. <i>European Journal of Medicinal Chemistry</i> , 2005, 40, 1134-1142.	2.6	32
9	Synthesis, SAR, and Biological Evaluation of ±-Sulfonylphosphonic Acids as Selective Matrix Metalloproteinase Inhibitors. <i>ChemMedChem</i> , 2009, 4, 352-362.	1.6	31
10	Catechol-based matrix metalloproteinase inhibitors with additional antioxidative activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 25-37.	2.5	29
11	Lactoferrin-derived Peptides Active towards Influenza: Identification of Three Potent Tetrapeptide Inhibitors. <i>Scientific Reports</i> , 2017, 7, 10593.	1.6	28
12	Synthesis and evaluation of new tripeptide phosphonate inhibitors of MMP-8 and MMP-2. <i>European Journal of Medicinal Chemistry</i> , 2005, 40, 271-279.	2.6	23
13	Fragmenting the S100B-p53 Interaction: Combined Virtual/Biophysical Screening Approaches to Identify Ligands. <i>ChemMedChem</i> , 2010, 5, 428-435.	1.6	22
14	Probing the S100B Zinc-Binding Site for the Identification of Non-Zinc-Binding MMP-2 Inhibitors. <i>ChemMedChem</i> , 2013, 8, 1475-1482.	1.6	22
15	Identification of new anti-Candida compounds by ligand-based pharmacophore virtual screening. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 1703-1706.	2.5	19
16	Novel bisphosphonates with antiresorptive effect in bone mineralization and osteoclastogenesis. <i>European Journal of Medicinal Chemistry</i> , 2018, 158, 184-200.	2.6	19
17	Seeking for Non-Zinc-Binding MMP-2 Inhibitors: Synthesis, Biological Evaluation and Molecular Modelling Studies. <i>International Journal of Molecular Sciences</i> , 2016, 17, 1768.	1.8	17
18	Development of CDK4/6 Inhibitors: A Five Years Update. <i>Molecules</i> , 2021, 26, 1488.	1.7	17

#	ARTICLE	IF	CITATIONS
19	Peptidyl 3-substituted 1-hydroxyureas as isosteric analogues of succinylhydroxamate MMP inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2008, 43, 1008-1014.	2.6	16
20	Fragment-Based Discovery of 5-Arylisatin-Based Inhibitors of Matrix Metalloproteinases 2 and 13. <i>ChemMedChem</i> , 2016, 11, 1892-1898.	1.6	16
21	PPAR α agonists based on stilbene and its bioisosteres: biological evaluation and docking studies. <i>MedChemComm</i> , 2015, 6, 1513-1517.	3.5	13
22	Amino Acid Derivatives as New Zinc Binding Groups for the Design of Selective Matrix Metalloproteinase Inhibitors. <i>Journal of Amino Acids</i> , 2013, 2013, 1-12.	5.8	12
23	Phosphonate Emerging Zinc Binding Group in Matrix Metalloproteinase Inhibitors. <i>Current Drug Targets</i> , 2015, 16, 1634-1644.	1.0	12
24	Non-Zinc-Binding Inhibitors of MMP-13: GRID-Based Approaches to Rationalize the Binding Process. <i>Current Topics in Medicinal Chemistry</i> , 2015, 16, 449-459.	1.0	11
25	An Effective Virtual Screening Protocol To Identify Promising p53-MDM2 Inhibitors. <i>Journal of Chemical Information and Modeling</i> , 2016, 56, 1216-1227.	2.5	10
26	Virtual screening identification and chemical optimization of substituted 2-arylbenzimidazoles as new non-zinc-binding MMP-2 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2020, 28, 115257.	1.4	10
27	In vitro comparison of new bisphosphonic acids and zoledronate effects on human gingival fibroblasts viability, inflammation and matrix turnover. <i>Clinical Oral Investigations</i> , 2016, 20, 2013-2021.	1.4	9
28	Bisphosphonate matrix metalloproteinase inhibitors for the treatment of periodontitis: An in vitro study. <i>International Journal of Molecular Medicine</i> , 2018, 42, 651-657.	1.8	8
29	Investigation of the N-BP Binding at FPPS by Combined Computational Approaches. <i>Medicinal Chemistry</i> , 2015, 11, 417-431.	0.7	8
30	Identification of small molecules acting against H1N1 influenza A virus. <i>Virology</i> , 2016, 488, 249-258.	1.1	7
31	An Integrated Computational Approach to Rationalize the Activity of Non-Zinc-Binding MMP-2 Inhibitors. <i>PLoS ONE</i> , 2012, 7, e47774.	1.1	7
32	Discovery of 7-aminophenanthridin-6-one as a new scaffold for matrix metalloproteinase inhibitors with multitarget neuroprotective activity. <i>European Journal of Medicinal Chemistry</i> , 2021, 210, 113061.	2.6	6
33	Dual targeting of cancer-related human matrix metalloproteinases and carbonic anhydrases by chiral N-(biarylsulfonyl)-phosphonic acids. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017, 32, 1260-1264.	2.5	4
34	Discovery of a Novel Tetrapeptide against Influenza A Virus: Rational Design, Synthesis, Bioactivity Evaluation and Computational Studies. <i>Pharmaceuticals</i> , 2021, 14, 959.	1.7	4
35	Ac-tLeu-Asp-H is the minimal and highly effective human caspase-3 inhibitor: biological and in silico studies. <i>Amino Acids</i> , 2015, 47, 153-162.	1.2	3
36	Mimic catechins to develop selective MMP-2 inhibitors. <i>Monatshefte für Chemie</i> , 2018, 149, 1293-1300.	0.9	3

#	ARTICLE	IF	CITATIONS
37	Effects of Biphenyl Sulfonylamino Methyl Bisphosphonic Acids on Porphyromonas Gingivalis and Cytokine Secretion by Oral Epithelial Cells. Medicinal Chemistry, 2013, 9, 855-860.	0.7	3
38	Broad-Spectrum Activity of Small Molecules Acting against Influenza a Virus: Biological and Computational Studies. Pharmaceuticals, 2022, 15, 301.	1.7	3
39	Bone-Seeking Matrix Metalloproteinase Inhibitors for the Treatment of Skeletal Malignancy. Pharmaceuticals, 2020, 13, 113.	1.7	2
40	Het(aryl)isatin to het(aryl)aminoindoline scaffold hopping: A route to selective inhibitors of matrix metalloproteinases. Arabian Journal of Chemistry, 2022, 15, 103492.	2.3	2
41	(2-Aminobenzothiazole)-Methyl-1,1-Bisphosphonic Acids: Targeting Matrix Metalloproteinase 13 Inhibition to the Bone. Pharmaceuticals, 2021, 14, 85.	1.7	1