

Mariangela Agamennone

List of Publications by Year in descending order

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| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | Het(aryl)isatin to het(aryl)aminoindoline scaffold hopping: A route to selective inhibitors of matrix metalloproteinases. <i>Arabian Journal of Chemistry</i> , 2022, 15, 103492. | 2.3 | 2 |
| 2 | Broad-Spectrum Activity of Small Molecules Acting against Influenza a Virus: Biological and Computational Studies. <i>Pharmaceuticals</i> , 2022, 15, 301. | 1.7 | 3 |
| 3 | 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 |
| 4 | (2-Aminobenzothiazole)-Methyl-1,1-Bisphosphonic Acids: Targeting Matrix Metalloproteinase 13 Inhibition to the Bone. <i>Pharmaceuticals</i> , 2021, 14, 85. | 1.7 | 1 |
| 5 | Development of CDK4/6 Inhibitors: A Five Years Update. <i>Molecules</i> , 2021, 26, 1488. | 1.7 | 17 |
| 6 | 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 |
| 7 | 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 |
| 8 | Bone-Seeking Matrix Metalloproteinase Inhibitors for the Treatment of Skeletal Malignancy. <i>Pharmaceuticals</i> , 2020, 13, 113. | 1.7 | 2 |
| 9 | 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 |
| 10 | 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 |
| 11 | Novel bisphosphonates with antiresorptive effect in bone mineralization and osteoclastogenesis. <i>European Journal of Medicinal Chemistry</i> , 2018, 158, 184-200. | 2.6 | 19 |
| 12 | Mimic catechins to develop selective MMP-2 inhibitors. <i>Monatshefte für Chemie</i> , 2018, 149, 1293-1300. | 0.9 | 3 |
| 13 | Dual targeting of cancer-related human matrix metalloproteinases and carbonic anhydrases by chiral <i>N</i> -(biarylsulfonyl)-phosphonic acids. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017, 32, 1260-1264. | 2.5 | 4 |
| 14 | Lactoferrin-derived Peptides Active towards Influenza: Identification of Three Potent Tetrapeptide Inhibitors. <i>Scientific Reports</i> , 2017, 7, 10593. | 1.6 | 28 |
| 15 | 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 |
| 16 | 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 |
| 17 | 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 |
| 18 | Fragment-Based Discovery of Arylisatin-Based Inhibitors of Matrix Metalloproteinases 2 and 13. <i>ChemMedChem</i> , 2016, 11, 1892-1898. | 1.6 | 16 |

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|----|---|-----|-----------|
| 19 | 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 |
| 20 | Identification of new anti- <i>Candida</i> compounds by ligand-based pharmacophore virtual screening. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 1703-1706. | 2.5 | 19 |
| 21 | Identification of small molecules acting against H1N1 influenza A virus. <i>Virology</i> , 2016, 488, 249-258. | 1.1 | 7 |
| 22 | 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 |
| 23 | PPAR α agonists based on stilbene and its bioisosteres: biological evaluation and docking studies. <i>MedChemComm</i> , 2015, 6, 1513-1517. | 3.5 | 13 |
| 24 | Phosphonate Emerging Zinc Binding Group in Matrix Metalloproteinase Inhibitors. <i>Current Drug Targets</i> , 2015, 16, 1634-1644. | 1.0 | 12 |
| 25 | 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 |
| 26 | Investigation of the N-BP Binding at FPPS by Combined Computational Approaches. <i>Medicinal Chemistry</i> , 2015, 11, 417-431. | 0.7 | 8 |
| 27 | Probing the S1 α Site for the Identification of Non-Zinc-Binding MMP Inhibitors. <i>ChemMedChem</i> , 2013, 8, 1475-1482. | 1.6 | 22 |
| 28 | Arylamino methylene bisphosphonate derivatives as bone seeking matrix metalloproteinase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 6456-6465. | 1.4 | 34 |
| 29 | 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 |
| 30 | Effects of Biphenyl Sulfonylamino Methyl Bisphosphonic Acids on <i>Porphyromonas Gingivalis</i> and Cytokine Secretion by Oral Epithelial Cells. <i>Medicinal Chemistry</i> , 2013, 9, 855-860. | 0.7 | 3 |
| 31 | 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 |
| 32 | 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 |
| 33 | Biphenyl Sulfonylamino Methyl Bisphosphonic Acids as Inhibitors of Matrix Metalloproteinases and Bone Resorption. <i>ChemMedChem</i> , 2011, 6, 1258-1268. | 1.6 | 44 |
| 34 | Fragmenting the S100B-p53 Interaction: Combined Virtual/Biophysical Screening Approaches to Identify Ligands. <i>ChemMedChem</i> , 2010, 5, 428-435. | 1.6 | 22 |
| 35 | Synthesis, SAR, and Biological Evaluation of α -Sulfonylphosphonic Acids as Selective Matrix Metalloproteinase Inhibitors. <i>ChemMedChem</i> , 2009, 4, 352-362. | 1.6 | 31 |
| 36 | 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 |

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|----|--|-----|-----------|
| 37 | Î±-Biphenylsulfonylamino 2-methylpropyl phosphonates: Enantioselective synthesis and selective inhibition of MMPs. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 791-799. | 1.4 | 39 |
| 38 | 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 |
| 39 | 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 |
| 40 | 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 |
| 41 | AMBER force field implementation of the boronate function to simulate the inhibition of Î²-lactamases by alkyl and aryl boronic acids. <i>European Journal of Medicinal Chemistry</i> , 2005, 40, 1134-1142. | 2.6 | 32 |