## Elisa Magli

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	5-HT <sub>1A</sub> Receptor: An Old Target as a New Attractive Tool in Drug Discovery from Central Nervous System to Cancer. Journal of Medicinal Chemistry, 2014, 57, 4407-4426.	2.9	85
2	Level, potential sources of polycyclic aromatic hydrocarbons (PAHs) in particulate matter (PM10) in Naples. Atmospheric Environment, 2016, 129, 186-196.	1.9	45
3	1,2,4-Thiadiazolidin-3,5-diones as novel hydrogen sulfide donors. European Journal of Medicinal Chemistry, 2018, 143, 1677-1686.	2.6	38
4	Trends in H2S-Donors Chemistry and Their Effects in Cardiovascular Diseases. Antioxidants, 2021, 10, 429.	2.2	38
5	Heavy Metals Size Distribution in PM10 and Environmental-Sanitary Risk Analysis in Acerra (Italy). Atmosphere, 2018, 9, 58.	1.0	37
6	H2S Donors and Their Use in Medicinal Chemistry. Biomolecules, 2021, 11, 1899.	1.8	36
7	Structure-activity relationships study of isothiocyanates for H2S releasing properties: 3-Pyridyl-isothiocyanate as a new promising cardioprotective agent. Journal of Advanced Research, 2021, 27, 41-53.	4.4	28
8	5-HT 2 receptor affinity, docking studies and pharmacological evaluation of a series of 1,3-disubstituted thiourea derivatives. European Journal of Medicinal Chemistry, 2016, 116, 173-186.	2.6	23
9	Anti-metastatic Properties of Naproxen-HBTA in a Murine Model of Cutaneous Melanoma. Frontiers in Pharmacology, 2019, 10, 66.	1.6	22
10	Fragment-based de novo design of a cystathionine Î <sup>3</sup> -lyase selective inhibitor blocking hydrogen sulfide production. Scientific Reports, 2016, 6, 34398.	1.6	20
11	Efficient microwave combinatorial synthesis of novel indolic arylpiperazine derivatives as serotoninergic ligands. European Journal of Medicinal Chemistry, 2010, 45, 752-759.	2.6	19
12	Microwave Assisted Organic Synthesis of Heterocycles in Aqueous Media: Recent Advances in Medicinal Chemistry. Medicinal Chemistry, 2016, 12, 720-732.	0.7	19
13	The Role of 5-HT1A Receptor in Cancer as a New Opportunity in Medicinal Chemistry. Current Medicinal Chemistry, 2018, 25, 3214-3227.	1.2	18
14	New 5-HT1A receptor ligands containing a N′-cyanoisonicotinamidine nucleus: Synthesis and in vitro pharmacological evaluation. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 2978-2982.	1.0	17
15	New 5-HT1A, 5HT2A and 5HT2C receptor ligands containing a picolinic nucleus: Synthesis, in vitro and in vivo pharmacological evaluation. Bioorganic and Medicinal Chemistry, 2017, 25, 5820-5837.	1.4	17
16	H2S donating corticosteroids: Design, synthesis and biological evaluation in a murine model of asthma. Journal of Advanced Research, 2022, 35, 267-277.	4.4	17
17	Design, synthesis and biological evaluation of TAR and cTAR binders as HIV-1 nucleocapsid inhibitors. MedChemComm, 2013, 4, 1388.	3.5	16
18	Synthesis and in Vitro Screening of New Series of 2,6-Dipeptidyl-anthraquinones: Influence of Side Chain Length on HIV-1 Nucleocapsid Inhibitors. Journal of Medicinal Chemistry, 2016, 59, 1914-1924.	2.9	15

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19	Synthesis, inÂvitro and inÂvivo pharmacological evaluation of serotoninergic ligands containing an isonicotinic nucleus. European Journal of Medicinal Chemistry, 2016, 110, 133-150.	2.6	14
20	New potent 5-HT2A receptor ligands containing an N′-cyanopicolinamidine nucleus: Synthesis and inÂvitro pharmacological evaluation. European Journal of Medicinal Chemistry, 2012, 47, 520-529.	2.6	12
21	Synthesis of 1-naphtylpiperazine derivatives as serotoninergic ligands and their evaluation as antiproliferative agents. European Journal of Medicinal Chemistry, 2011, 46, 2206-2216.	2.6	11
22	Chemical Composition of PM10 at Urban Sites in Naples (Italy). Atmosphere, 2016, 7, 163.	1.0	11
23	Synthesis of benzamide derivatives and their evaluation as antiprion agents. Bioorganic and Medicinal Chemistry, 2012, 20, 5001-5011.	1.4	10
24	Hybrids between H2S-donors and betamethasone 17-valerate or triamcinolone acetonide inhibit mast cell degranulation and promote hyperpolarization of bronchial smooth muscle cells. European Journal of Medicinal Chemistry, 2021, 221, 113517.	2.6	10
25	Synthesis and <i>In Vitro</i> Pharmacological Evaluation of Novel 2â€Hydroxypropylâ€4â€arylpiperazine Derivatives as Serotoninergic Ligands. Archiv Der Pharmazie, 2014, 347, 698-706.	2.1	9
26	New arylpiperazine derivatives with antidepressant-like activity containing isonicotinic and picolinic nuclei: evidence for serotonergic system involvement. Naunyn-Schmiedeberg's Archives of Pharmacology, 2019, 392, 743-754.	1.4	9
27	Design of Sphingosine Kinases Inhibitors: Challenges and Recent Developments. Current Pharmaceutical Design, 2019, 25, 956-968.	0.9	9
28	Prolonged NCX activation prevents SOD1 accumulation, reduces neuroinflammation, ameliorates motor behavior and prolongs survival in a ALS mouse model. Neurobiology of Disease, 2021, 159, 105480.	2.1	8
29	Non-Natural Linker Configuration in 2,6-Dipeptidyl-Anthraquinones Enhances the Inhibition of TAR RNA Binding/Annealing Activities by HIV-1 NC and Tat Proteins. Bioconjugate Chemistry, 2018, 29, 2195-2207.	1.8	7
30	Synthesis, docking studies, and pharmacological evaluation of 5HT <sub>2C</sub> ligands containing the <i>N</i> ′â€ɛyanoisonicotinamidine or <i>N</i> ′â€ɛyanopicolinamidine nucleus. Archiv Der Pharmazie, 2019, 352, e1800373.	2.1	7
31	PCB levels in adipose tissue of dogs from illegal dumping sites in Campania region (Italy). Chemosphere, 2020, 244, 125478.	4.2	7
32	Synthesis, docking studies, and pharmacological evaluation of 2â€hydroxypropylâ€4â€arylpiperazine derivatives as serotoninergic ligands. Archiv Der Pharmazie, 2021, 354, 2000414.	2.1	7
33	New Insights into the Structure–Activity Relationship and Neuroprotective Profile of Benzodiazepinone Derivatives of <b>Neurounina-1</b> as Modulators of the Na <sup>+</sup> /Ca <sup>2+</sup> Exchanger Isoforms. Journal of Medicinal Chemistry, 2021, 64, 17901-17919.	2.9	6
34	Development of 1,2,3-Triazole-Based Sphingosine Kinase Inhibitors and Their Evaluation as Antiproliferative Agents. International Journal of Molecular Sciences, 2017, 18, 2332.	1.8	5
35	Development, Validation of LC-MS/MS Method and Determination of Pharmacokinetic Parameters of the Stroke Neuroprotectant Neurounina-1 in Beagle Dog Plasma After Intravenous Administration. Frontiers in Pharmacology, 2019, 10, 432.	1.6	5
36	Synthesis, in vitro and in vivo evaluation of 11C-O-methylated arylpiperazines as potential serotonin 1A (5-HT1A) receptor antagonist radiotracers. EJNMMI Radiopharmacy and Chemistry, 2020, 5, 13.	1.8	5

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37	Serotoninergic receptor ligands improve Tamoxifen effectiveness on breast cancer cells. BMC Cancer, 2022, 22, 171.	1.1	4
38	Synthesis and Pharmacological Screening of Pyridopyrimidines as Effective Antiâ€Diarrheal Agents through the Suppression of Cyclic Nucleotide Accumulation. ChemistryOpen, 2019, 8, 464-475.	0.9	3
39	Multiple <i>in Vitro</i> Inhibition of HIV-1 Proteins by 2,6-Dipeptidyl-anthraquinone Conjugates Targeting the PBS RNA. ACS Medicinal Chemistry Letters, 2020, 11, 949-955.	1.3	1
40	Antagonizing S1P3 Receptor with Cell-Penetrating Pepducins in Skeletal Muscle Fibrosis. International Journal of Molecular Sciences, 2021, 22, 8861.	1.8	1
41	New Serotoninergic Ligands Containing Indolic and Methyl Indolic Nuclei: Synthesis and In Vitro Pharmacological Evaluation. Medicinal Chemistry, 2020, 16, 517-530.	0.7	1
42	Synthesis of Arylpiperazine Derivatives as Protease Activated Receptor 1 Antagonists and Their Evaluation as Antiproliferative Agents. Anti-Cancer Agents in Medicinal Chemistry, 2017, 17, 973-981.	0.9	0
43	Propafenone quantification in human plasma by high-performance liquid chromatography coupled with electrospray tandem mass spectrometry in a bioequivalence study. International Journal of	0.3	Ο