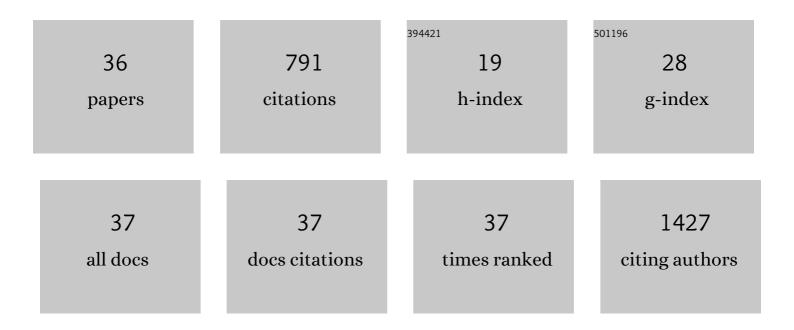
Marina Sala

List of Publications by Year in descending order

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MADINIA SALA

#	Article	IF	CITATIONS
1	Structure-Based Development of SARS-CoV-2 Spike Interactors. International Journal of Molecular Sciences, 2022, 23, 5601.	4.1	3
2	Discovery of a Novel Tetrapeptide against Influenza A Virus: Rational Design, Synthesis, Bioactivity Evaluation and Computational Studies. Pharmaceuticals, 2021, 14, 959.	3.8	4
3	Identification of a dual acting SARS-CoV-2 proteases inhibitor through in silico design and step-by-step biological characterization. European Journal of Medicinal Chemistry, 2021, 226, 113863.	5.5	22
4	Overcome Chemoresistance: Biophysical and Structural Analysis of Synthetic FHIT-Derived Peptides. Frontiers in Molecular Biosciences, 2021, 8, 715263.	3.5	1
5	Exploiting GRK2 Inhibition as a Therapeutic Option in Experimental Cancer Treatment: Role of p53-Induced Mitochondrial Apoptosis. Cancers, 2020, 12, 3530.	3.7	6
6	Role of Endothelial G Protein-Coupled Receptor Kinase 2 in Angioedema. Hypertension, 2020, 76, 1625-1636.	2.7	23
7	Synthesis and Ex Vivo Trans-Corneal Permeation of Penetratin Analogues as Ophthalmic Carriers: Preliminary Results. Pharmaceutics, 2020, 12, 728.	4.5	4
8	Pharmacological inhibition of <scp>GRK2</scp> improves cardiac metabolism and function in experimental heart failure. ESC Heart Failure, 2020, 7, 1571-1584.	3.1	21
9	In silico Analysis Revealed Potential Anti-SARS-CoV-2 Main Protease Activity by the Zonulin Inhibitor Larazotide Acetate. Frontiers in Chemistry, 2020, 8, 628609.	3.6	21
10	Antidiabetic and Cardioprotective Effects of Pharmacological Inhibition of GRK2 in db/db Mice. International Journal of Molecular Sciences, 2019, 20, 1492.	4.1	22
11	Identification of the 2-Benzoxazol-2-yl-phenol Scaffold as New Hit for JMJD3 Inhibition. ACS Medicinal Chemistry Letters, 2019, 10, 601-605.	2.8	23
12	<i>⁽ⁱ)^Î²</i> -Lactoglobulin Heptapeptide Reduces Oxidative Stress in Intestinal Epithelial Cells and Angiotensin II-Induced Vasoconstriction on Mouse Mesenteric Arteries by Induction of Nuclear Factor Erythroid 2-Related Factor 2 (Nrf2) Translocation. Oxidative Medicine and Cellular Longevity, 2019, 2019, 1-13.	4.0	12
13	Novel Potent Decameric Peptide of <i>Spirulina platensis</i> Reduces Blood Pressure Levels Through a PI3K/AKT/eNOS-Dependent Mechanism. Hypertension, 2019, 73, 449-457.	2.7	53
14	Chemical profiling of bioactive constituents in hop cones and pellets extracts by online comprehensive twoâ€dimensional liquid chromatography with tandem mass spectrometry and direct infusion Fourier transform ion cyclotron resonance mass spectrometry. Journal of Separation Science, 2018, 41, 1548-1557.	2.5	36
15	A Novel Small Peptide Inhibitor of NFκB, RH10, Blocks Oxidative Stress-Dependent Phenotypes in Cancer. Oxidative Medicine and Cellular Longevity, 2018, 2018, 1-9.	4.0	4
16	Identification of a Potent Tryptophan-Based TRPM8 Antagonist With in Vivo Analgesic Activity. Journal of Medicinal Chemistry, 2018, 61, 6140-6152.	6.4	31
17	Investigation on side-product formation during the synthesis of a lactoferrin-derived lactam-bridged cyclic peptide. Amino Acids, 2018, 50, 1367-1375.	2.7	1
18	Antioxidant Properties of Buffalo-Milk Dairy Products: A β-Lg Peptide Released after Gastrointestinal Digestion of Buffalo Ricotta Cheese Reduces Oxidative Stress in Intestinal Epithelial Cells. International Journal of Molecular Sciences, 2018, 19, 1955.	4.1	42

MARINA SALA

#	Article	IF	CITATIONS
19	Design, Synthesis, Biological Activity, and Structural Analysis of Lactamâ€Constrained PTPRJ Agonist Peptides. ChemMedChem, 2018, 13, 1673-1680.	3.2	1
20	Cell penetrating peptides in ocular drug delivery: State of the art. Journal of Controlled Release, 2018, 284, 84-102.	9.9	84
21	Synergistic effect of divalent cations in improving technological properties of cross-linked alginate beads. International Journal of Biological Macromolecules, 2017, 101, 100-106.	7.5	38
22	On the design of tailored liposomes for KRX29 peptide delivery. New Journal of Chemistry, 2017, 41, 11280-11290.	2.8	2
23	Lactoferrin-derived Peptides Active towards Influenza: Identification of Three Potent Tetrapeptide Inhibitors. Scientific Reports, 2017, 7, 10593.	3.3	28
24	Bioavailable Citrus sinensis Extract: Polyphenolic Composition and Biological Activity. Molecules, 2017, 22, 623.	3.8	31
25	Anti-Inflammatory, Antioxidant and Crystallographic Studies of N-Palmitoyl-ethanol Amine (PEA) Derivatives. Molecules, 2017, 22, 616.	3.8	15
26	Development and Identification of a Novel Anti-HIV-1 Peptide Derived by Modification of the N-Terminal Domain of HIV-1 Integrase. Frontiers in Microbiology, 2016, 7, 845.	3.5	13
27	Design and Synthesis of New Cell Penetrating Peptides: Diffusion and Distribution Inside the Cornea. Molecular Pharmaceutics, 2016, 13, 3876-3883.	4.6	24
28	Identification of an indol-based derivative as potent and selective varicella zoster virus (VZV) inhibitor. European Journal of Medicinal Chemistry, 2016, 124, 773-781.	5.5	15
29	Targeting the CaMKII/ERK Interaction in the Heart Prevents Cardiac Hypertrophy. PLoS ONE, 2015, 10, e0130477.	2.5	52
30	SAR study and conformational analysis of a series of novel peptide G protein oupled receptor kinase 2 inhibitors. Biopolymers, 2014, 101, 121-128.	2.4	10
31	Discovery of PTPRJ Agonist Peptides That Effectively Inhibit <i>in Vitro</i> Cancer Cell Proliferation and Tube Formation. ACS Chemical Biology, 2013, 8, 1497-1506.	3.4	27
32	Design, synthesis and efficacy of novel G protein-coupled receptor kinase 2 inhibitors. European Journal of Medicinal Chemistry, 2013, 69, 384-392.	5.5	19
33	Biological activity of 3-chloro-azetidin-2-one derivatives having interesting antiproliferative activity on human breast cancer cell lines. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 6401-6405.	2.2	45
34	Characterization of a selective CaMKII peptide inhibitor. European Journal of Medicinal Chemistry, 2013, 62, 425-434.	5.5	22
35	Design, Synthesis, and Evaluation of New Tripeptides as COX-2 Inhibitors. Journal of Amino Acids, 2013, 2013, 1-7.	5.8	4
36	Isolation and Functional Characterization of Peptide Agonists of PTPRJ, a Tyrosine Phosphatase Receptor Endowed with Tumor Suppressor Activity. ACS Chemical Biology, 2012, 7, 1666-1676.	3.4	32