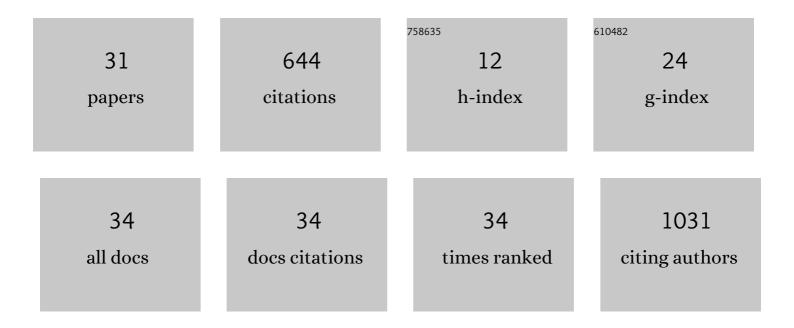
Nuria Cirauqui

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Using normal mode analysis on protein structural models. How far can we go on our predictions?. Proteins: Structure, Function and Bioinformatics, 2021, 89, 531-543.	1.5	16
2	Three-Finger Toxins from Brazilian Coral Snakes: From Molecular Framework to Insights in Biological Function. Toxins, 2021, 13, 328.	1.5	3
3	Structural insights into the allosteric site of Arabidopsis NADP-malic enzyme 2: role of the second sphere residues in the regulatory signal transmission. Plant Molecular Biology, 2021, 107, 37-48.	2.0	1
4	Diterpenes isolated from <i>Canistrocarpus cervicornis</i> with virucidal activity against HIV-1: an <i>in silico</i> evaluation. Natural Product Research, 2021, , 1-5.	1.0	1
5	In Vivo Biological Evaluation of a Synthetic Royleanone Derivative as a Promising Fast-Acting Trypanocidal Agent by Inducing Mitochondrial-Dependent Necrosis. Journal of Natural Products, 2020, 83, 3571-3583.	1.5	6
6	Aerolysin nanopores decode digital information stored in tailored macromolecular analytes. Science Advances, 2020, 6, .	4.7	57
7	Phase Separation and Disorder-to-Order Transition of Human Brain Expressed X-Linked 3 (hBEX3) in the Presence of Small Fragments of tRNA. Journal of Molecular Biology, 2020, 432, 2319-2348.	2.0	13
8	Repositioning of leishmanicidal [1,2,3]Triazolo[1,5-a]pyridinium salts for Chagas disease treatment: Trypanosoma cruzi cell death involving mitochondrial membrane depolarisation and Fe-SOD inhibition. Parasitology Research, 2020, 119, 2943-2954.	0.6	4
9	Rational modification of Mannich base-type derivatives as novel antichagasic compounds: Synthesis, in vitro and in vivo evaluation. Bioorganic and Medicinal Chemistry, 2019, 27, 3902-3917.	1.4	17
10	Antichagasic profile of a Series of Mannich Base‶ype Derivatives: Design, Synthesis, <i>inâ€vitro</i> Evaluation, and Computational Studies Involving Iron Superoxide Dismutase. ChemistrySelect, 2019, 4, 8112-8121.	0.7	3
11	Single-molecule sensing of peptides and nucleic acids by engineered aerolysin nanopores. Nature Communications, 2019, 10, 4918.	5.8	74
12	A step towards development of promising trypanocidal agents: Synthesis, characterization and inÂvitro biological evaluation of ferrocenyl Mannich base-type derivatives. European Journal of Medicinal Chemistry, 2019, 163, 569-582.	2.6	11
13	BK polyomavirus genotypes la and lb1 exhibit different biological properties in renal transplant recipients. Virus Research, 2018, 243, 65-68.	1.1	17
14	Mapping the sensing spots of aerolysin for single oligonucleotides analysis. Nature Communications, 2018, 9, 2823.	5.8	60
15	Second Generation of Mannich Base-Type Derivatives with <i>in Vivo</i> Activity against <i>Trypanosoma cruzi</i> . Journal of Medicinal Chemistry, 2018, 61, 5643-5663.	2.9	32
16	Structural, physicochemical and dynamic features conserved within the aerolysin pore-forming toxin family. Scientific Reports, 2017, 7, 13932.	1.6	38
17	Analysis of worldwide sequence mutations in Zika virus proteins E, NS1, NS3 and NS5 from a structural point of view. Molecular BioSystems, 2017, 13, 122-131.	2.9	8
18	Biological Evaluation of Arylamine Mannich Base Derivatives with Potent In Vivo Activity as Potent Antichagasic Agents. Proceedings (mdpi), 2017, 1, .	0.2	0

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19	Cryo-EM structure of aerolysin variants reveals a novel protein fold and the pore-formation process. Nature Communications, 2016, 7, 12062.	5.8	144
20	Phylogenetic and structural analysis of merkel cell polyomavirus VP1 in Brazilian samples. Virus Research, 2016, 221, 1-7.	1.1	8
21	In Vitro and in Vivo Anti-Trypanosoma cruziActivity of New Arylamine Mannich Base-Type Derivatives. Journal of Medicinal Chemistry, 2016, 59, 10929-10945.	2.9	30
22	Design, Synthesis and Evaluation of New Fluoroamodiaquine Analogues. Chemical and Pharmaceutical Bulletin, 2016, 64, 594-601.	0.6	6
23	Genetic and structural analysis of polyomavirus BK Tâ€antigens reveal a higher density of mutations at interâ€domain and hexamerization regions, regardless the status of infection. Journal of Medical Virology, 2015, 87, 1418-1426.	2.5	3
24	Molecular modeling study of a series of amodiaquine analogues with antimalarial activity. Medicinal Chemistry Research, 2015, 24, 3529-3536.	1.1	5
25	Genetic and Structural Analysis of Merkel Cell Polyomavirus Large T Antigen from Diverse Biological Samples. Intervirology, 2014, 57, 331-336.	1.2	1
26	Novel sulfonylurea derivatives as H3 receptor antagonists. Preliminary SAR studies. European Journal of Medicinal Chemistry, 2012, 52, 1-13.	2.6	15
27	Building a MCHR1 homology model provides insight into the receptor–antagonist contacts that are important for the development of new anti-obesity agents. Bioorganic and Medicinal Chemistry, 2010, 18, 7365-7379.	1.4	8
28	New Amide Derivatives as Melanin-concentrating Hormone Receptor 1 Antagonists for the Treatment of Obesity. Arzneimittelforschung, 2008, 58, 585-591.	0.5	0
29	Melanin-Concentrating Hormone Receptor 1 Antagonists: A New Perspective for the Pharmacologic Treatment of Obesity. Current Medicinal Chemistry, 2008, 15, 1025-1043.	1.2	50
30	Novel series of substituted biphenylmethyl urea derivatives as MCH-R1 antagonists for the treatment of obesity. Bioorganic and Medicinal Chemistry, 2007, 15, 3896-3911.	1.4	11
31	Multiscale molecular simulations to investigate adenylyl cyclaseâ€based signaling in the brain. Wiley Interdisciplinary Reviews: Computational Molecular Science, 0, , .	6.2	2