Isabella Alvim Guedes

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7066621/publications.pdf

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21 papers

1,097 citations

759233 12 h-index 17 g-index

22 all docs 22 docs citations

times ranked

22

1494 citing authors

#	Article	IF	CITATIONS
1	Isobenzofuran-1(3H)-ones as new tyrosinase inhibitors: Biological activity and interaction studies by molecular docking and NMR. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2021, 1869, 140580.	2.3	6
2	New machine learning and physics-based scoring functions for drug discovery. Scientific Reports, 2021, 11, 3198.	3.3	91
3	Drug design and repurposing with DockThor-VS web server focusing on SARS-CoV-2 therapeutic targets and their non-synonym variants. Scientific Reports, 2021, 11, 5543.	3.3	63
4	Highly Flexible Ligand Docking: Benchmarking of the DockThor Program on the LEADS-PEP Protein–Peptide Data Set. Journal of Chemical Information and Modeling, 2020, 60, 667-683.	5.4	144
5	Cinnamoyl-N-Acylhydrazone-Donepezil Hybrids: Synthesis and Evaluation of Novel Multifunctional Ligands Against Neurodegenerative Diseases. Neurochemical Research, 2020, 45, 3003-3020.	3.3	7
6	Design, Synthesis and Biological Evaluation of Novel Triazole N-acylhydrazone Hybrids for Alzheimer's Disease. Molecules, 2020, 25, 3165.	3.8	14
7	Synthesis of new lophine–carbohydrate hybrids as cholinesterase inhibitors: cytotoxicity evaluation and molecular modeling. MedChemComm, 2019, 10, 2089-2101.	3.4	13
8	Design, synthesis and pharmacological evaluation of N -benzyl-piperidinyl-aryl-acylhydrazone derivatives as donepezil hybrids: Discovery of novel multi-target anti-alzheimer prototype drug candidates. European Journal of Medicinal Chemistry, 2018, 147, 48-65.	5 . 5	52
9	Discovery of naphthylâ€ <i>N</i> àâ€acylhydrazone p38α MAPK inhibitors with in vivo antiâ€inflammatory and antiâ€TNFâ€Î± activity. Chemical Biology and Drug Design, 2018, 91, 391-397.	3.2	22
10	Empirical Scoring Functions for Structure-Based Virtual Screening: Applications, Critical Aspects, and Challenges. Frontiers in Pharmacology, 2018, 9, 1089.	3. 5	185
11	Design, synthesis, cholinesterase inhibition and molecular modelling study of novel tacrine hybrids with carbohydrate derivatives. Bioorganic and Medicinal Chemistry, 2018, 26, 5566-5577.	3.0	21
12	Design, synthesis and evaluation of novel feruloyl-donepezil hybrids as potential multitarget drugs for the treatment of Alzheimer's disease. European Journal of Medicinal Chemistry, 2017, 130, 440-457.	5 . 5	67
13	LASSBioâ€1829 Hydrochloride: Development of a New Orally Active <i>N</i> â€Acylhydrazone IKK2 Inhibitor with Antiâ€inflammatory Properties. ChemMedChem, 2016, 11, 234-244.	3.2	7
14	Novel series of tacrine-tianeptine hybrids: Synthesis, cholinesterase inhibitory activity, \$100B secretion and a molecular modeling approach. European Journal of Medicinal Chemistry, 2016, 121, 758-772.	5 . 5	39
15	A unique SaeS allele overrides cell-density dependent expression of saeR and lukSF-PV in the ST30-SCCmecIV lineage of CA-MRSA. International Journal of Medical Microbiology, 2016, 306, 367-380.	3.6	10
16	Structural modeling and docking studies of ribose 5-phosphate isomerase from Leishmania major and Homo sapiens: A comparative analysis for Leishmaniasis treatment. Journal of Molecular Graphics and Modelling, 2015, 55, 134-147.	2.4	23
17	Receptor–ligand molecular docking. Biophysical Reviews, 2014, 6, 75-87.	3.2	324
18	Chiral Bistacrine Analogues: Synthesis, Cholinesterase Inhibitory Activity and a Molecular Modeling Approach. Journal of the Brazilian Chemical Society, 0, , .	0.6	5

#	Article	IF	CITATIONS
19	Expedient Microwave-Assisted Synthesis of Bis(n)-lophine Analogues as Selective Butyrylcholinesterase Inhibitors: Cytotoxicity Evaluation and Molecular Modelling. Journal of the Brazilian Chemical Society, 0, , .	0.6	1
20	An Expedient Synthesis of Tacrine-Squaric Hybrids as Potent, Selective and Dual‑Binding Cholinesterase Inhibitors. Journal of the Brazilian Chemical Society, 0, , .	0.6	0
21	Design, synthesis, and biological evaluation of new thalidomide–donepezil hybrids as neuroprotective agents targeting cholinesterases and neuroinflammation. RSC Medicinal Chemistry, 0, , .	3.9	1