

Lidia Ciccone

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3718141/publications.pdf>

Version: 2024-02-01

35
papers

571
citations

623734

14
h-index

677142

22
g-index

36
all docs

36
docs citations

36
times ranked

726
citing authors

#	ARTICLE	IF	CITATIONS
1	Synthesis and Evaluation of Monoaryl Derivatives as Transthyretin Fibril Formation Inhibitors. <i>Pharmaceutical Chemistry Journal</i> , 2022, 56, 38-47.	0.8	2
2	Nature-Inspired O-Benzyl Oxime-Based Derivatives as New Dual-Acting Agents Targeting Aldose Reductase and Oxidative Stress. <i>Biomolecules</i> , 2022, 12, 448.	4.0	11
3	Antioxidant Quercetin 3-O-Glycosylated Plant Flavonols Contribute to Transthyretin Stabilization. <i>Crystals</i> , 2022, 12, 638.	2.2	3
4	MT9, a natural peptide from black mamba venom antagonizes the muscarinic type 2 receptor and reverses the M2R-agonist-induced relaxation in rat and human arteries. <i>Biomedicine and Pharmacotherapy</i> , 2022, 150, 113094.	5.6	3
5	Pterostilbene fluorescent probes as potential tools for targeting neurodegeneration in biological applications. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022, 37, 1812-1820.	5.2	0
6	Identification of histone deacetylase inhibitors with (arylidene)aminoxy scaffold active in uveal melanoma cell lines. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021, 36, 34-47.	5.2	11
7	Activation of carbonic anhydrases from human brain by amino alcohol oxime ethers: towards human carbonic anhydrase VII selective activators. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021, 36, 48-57.	5.2	12
8	Application of PROTAC strategy to TTR-A β 2 protein-protein interaction for the development of Alzheimer's disease drugs. <i>Neural Regeneration Research</i> , 2021, 16, 1554.	3.0	10
9	Neuroglobin and neuroprotection: the role of natural and synthetic compounds in neuroglobin pharmacological induction. <i>Neural Regeneration Research</i> , 2021, 16, 2353.	3.0	12
10	Exosite inhibition of ADAMTS-5 by a glycoconjugated arylsulfonamide. <i>Scientific Reports</i> , 2021, 11, 949.	3.3	14
11	Natural Marine and Terrestrial Compounds as Modulators of Matrix Metalloproteinases-2 (MMP-2) and MMP-9 in Alzheimer's Disease. <i>Pharmaceuticals</i> , 2021, 14, 86.	3.8	26
12	Physiological Metals Can Induce Conformational Changes in Transthyretin Structure: Neuroprotection or Misfolding Induction?. <i>Crystals</i> , 2021, 11, 354.	2.2	9
13	Carbonic Anhydrase Inhibitors and Epilepsy: State of the Art and Future Perspectives. <i>Molecules</i> , 2021, 26, 6380.	3.8	27
14	Multifunctional Small Molecules as Potential Anti-Alzheimer's Disease Agents. <i>Molecules</i> , 2021, 26, 6015.	3.8	7
15	Discovery of Dimeric Arylsulfonamides as Potent ADAM8 Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 1787-1793.	2.8	3
16	Discovery of a new ATP-citrate lyase (ACLY) inhibitor identified by a pharmacophore-based virtual screening study. <i>Journal of Biomolecular Structure and Dynamics</i> , 2021, 39, 3996-4004.	3.5	4
17	Monoaryl derivatives as transthyretin fibril formation inhibitors: Design, synthesis, biological evaluation and structural analysis. <i>Bioorganic and Medicinal Chemistry</i> , 2020, 28, 115673.	3.0	8
18	Natural compounds as inhibitors of transthyretin amyloidosis and neuroprotective agents: analysis of structural data for future drug design. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020, 35, 1145-1162.	5.2	35

#	ARTICLE	IF	CITATIONS
19	The Positive Side of the Alzheimer's Disease Amyloid Cross-Interactions: The Case of the A β 1-42 Peptide with Tau, TTR, CysC, and ApoA1. <i>Molecules</i> , 2020, 25, 2439.	3.8	37
20	Oxy-imino saccharidic derivatives as a new structural class of aldose reductase inhibitors endowed with anti-oxidant activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020, 35, 1194-1205.	5.2	5
21	A Soluble Metabolon Synthesizes the Isoprenoid Lipid Ubiquinone. <i>Cell Chemical Biology</i> , 2019, 26, 482-492.e7.	5.2	46
22	Development of Thioaryl-Based Matrix Metalloproteinase-12 Inhibitors with Alternative Zinc-Binding Groups: Synthesis, Potentiometric, NMR, and Crystallographic Studies. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 4421-4435.	6.4	34
23	Copper mediated amyloid- β binding to Transthyretin. <i>Scientific Reports</i> , 2018, 8, 13744.	3.3	26
24	Comparison of helical scan and standard rotation methods in single-crystal X-ray data collection strategies. <i>Journal of Synchrotron Radiation</i> , 2017, 24, 42-52.	2.4	27
25	Bifunctional Inhibitors as a New Tool To Reduce Cancer Cell Invasion by Impairing MMP-9 Homodimerization. <i>ACS Medicinal Chemistry Letters</i> , 2017, 8, 293-298.	2.8	13
26	N-(Aroyl)-N-(arylmethoxy)- β -alanines: Selective inhibitors of aldose reductase. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 3068-3076.	3.0	13
27	Targeting Different Transthyretin Binding Sites with Unusual Natural Compounds. <i>ChemMedChem</i> , 2016, 11, 1865-1874.	3.2	16
28	Synthesis and structural analysis of halogen substituted fibril formation inhibitors of Human Transthyretin (TTR). <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 40-51.	5.2	15
29	Sugar-Based Arylsulfonamide Carboxylates as Selective and Water-Soluble Matrix Metalloproteinase-12 Inhibitors. <i>ChemMedChem</i> , 2016, 11, 1626-1637.	3.2	36
30	Human TTR conformation altered by rhenium tris-carbonyl derivatives. <i>Journal of Structural Biology</i> , 2016, 195, 353-364.	2.8	17
31	A new crystal form of human transthyretin obtained with a curcumin derived ligand. <i>Journal of Structural Biology</i> , 2016, 194, 8-17.	2.8	18
32	X-ray crystal structure and activity of fluorenyl-based compounds as transthyretin fibrillogenesis inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 824-833.	5.2	10
33	Multicomponent mixtures for cryoprotection and ligand solubilization. <i>Biotechnology Reports (Amsterdam, Netherlands)</i> , 2015, 7, 120-127.	4.4	22
34	Synthesis and cyclooxygenase inhibitory properties of new naphthalene-methylsulfonamido, naphthalene-methylsulfonyl and tetrahydronaphthalen-methylsulfonamido compounds. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015, 30, 406-412.	5.2	6
35	Transthyretin complexes with curcumin and bromo-estradiol: evaluation of solubilizing multicomponent mixtures. <i>New Biotechnology</i> , 2015, 32, 54-64.	4.4	33