

C David Pessoa-Mahana

List of Publications by Year in descending order

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papers

405
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687220

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31
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630
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#	ARTICLE	IF	CITATIONS
1	New Pyridone-Based Derivatives as Cannabinoid Receptor Type 2 Agonists. <i>International Journal of Molecular Sciences</i> , 2021, 22, 11212.	1.8	10
2	Synthesis, in vitro evaluation and molecular docking of a new class of indolylpropyl benzamidopiperazines as dual AChE and SERT ligands for Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2020, 198, 112368.	2.6	18
3	Cannabidiol binding and negative allosteric modulation at the cannabinoid type 1 receptor in the presence of delta-9-tetrahydrocannabinol: An In Silico study. <i>PLoS ONE</i> , 2019, 14, e0220025.	1.1	45
4	Three-Dimensional Quantitative Structure-Activity Relationships (3D-QSAR) on a Series of Piperazine-Carboxamides Fatty Acid Amide Hydrolase (FAAH) Inhibitors as a Useful Tool for the Design of New Cannabinoid Ligands. <i>International Journal of Molecular Sciences</i> , 2019, 20, 2510.	1.8	11
5	Synthesis and biological evaluation of potential acetylcholinesterase inhibitors based on a benzoxazine core. <i>Archiv Der Pharmazie</i> , 2018, 351, e1800024.	2.1	8
6	Combined CoMFA and CoMSIA 3D-QSAR study of benzimidazole and benzothiophene derivatives with selective affinity for the CB2 cannabinoid receptor. <i>European Journal of Pharmaceutical Sciences</i> , 2017, 101, 1-10.	1.9	20
7	Effect of alpha lipoic acid on leukotriene A 4 hydrolase. <i>European Journal of Pharmacology</i> , 2017, 799, 41-47.	1.7	2
8	Synthesis and Docking of Novel Indolylpropyl Derivatives as New Polypharmacological Agents Displaying Affinity for 5-HT _{1A} /SERT. <i>Archiv Der Pharmazie</i> , 2017, 350, e1600271.	2.1	9
9	Extended N-Arylsulfonylindoles as 5-HT ₆ Receptor Antagonists: Design, Synthesis & Biological Evaluation. <i>Molecules</i> , 2016, 21, 1070.	1.7	12
10	Synthesis, binding assays, cytotoxic activity and docking studies of benzimidazole and benzothiophene derivatives with selective affinity for the CB2 cannabinoid receptor. <i>European Journal of Medicinal Chemistry</i> , 2016, 124, 17-35.	2.6	25
11	Design, Synthesis, Biological Evaluation and Binding Mode Modeling of Benzimidazole Derivatives Targeting the Cannabinoid Receptor Type 1. <i>Archiv Der Pharmazie</i> , 2015, 348, 81-88.	2.1	7
12	3D-QSAR/CoMFA-Based Structure-Affinity/Selectivity Relationships of Aminoalkylindoles in the Cannabinoid CB1 and CB2 Receptors. <i>Molecules</i> , 2014, 19, 2842-2861.	1.7	17
13	Synthesis, docking and pharmacological evaluation of novel homo- and hetero-bis 3-piperazinylpropylindole derivatives at SERT and 5-HT _{1A} receptor. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 7604-7611.	1.4	12
14	Synthesis, Docking Studies and Biological Evaluation of Benzo[b]thiophen-2-yl-3-(4-arylpiperazin-1-yl)-propan-1-one Derivatives on 5-HT _{1A} Serotonin Receptors. <i>Molecules</i> , 2012, 17, 1388-1407.	1.7	14
15	SYNTHESIS OF 1-BENZYL-3-[4-(ARYL-1-PIPERAZINYL) CARBONYL]-1H-INDOLES: NOVEL LIGANDS WITH POTENTIAL D4 DOPAMINERGIC ACTIVITY. <i>Journal of the Chilean Chemical Society</i> , 2011, 56, 866-869.	0.5	3
16	Evaluation of the Membrane Permeability (PAMPA and Skin) of Benzimidazoles with Potential Cannabinoid Activity and their Relation with the Biopharmaceutics Classification System (BCS). <i>AAPS PharmSciTech</i> , 2011, 12, 573-578.	1.5	15
17	Synthesis of a novel series of 4-arylpiperazinyl derivatives linked to a 2-(pyridin-3-yl)-1H-benzimidazole as new Delavirdine analogues. <i>Journal of the Brazilian Chemical Society</i> , 2010, 21, 63-70.	0.6	9
18	Solvent-free microwave-promoted Michael addition of aza-nucleophiles to benzo[b]thiophen-2-yl-2-propenone. <i>Arquivoc</i> , 2009, 2009, 316-325.	0.3	7

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19	Microwave-assisted synthesis and regioisomeric structural elucidation of novel benzimidazo[1,2-d][1,4]benzodiazepinone derivatives. <i>Arkivoc</i> , 2009, 2009, 131-140.	0.3	14
20	SYNTHESIS OF CONFORMATIONALLY RESTRICTED N-{4-[4-(4,7-DIMETHOXY-BENZO[b]THIOPHENE-2-CARBONYL)-1-PIPERAZINYL]-PHENYL}-ARYLCARBOXAMIDES POTENTIAL LIGANDS WITH 5-HT1A BINDING AFFINITY. <i>Journal of the Chilean Chemical Society</i> , 2009, 54, .	0.5	0
21	Computational modeling study of functional microdomains in cannabinoid receptor type 1. <i>Bioorganic and Medicinal Chemistry</i> , 2008, 16, 4378-4389.	1.4	23
22	Solvent-Free Microwave Synthesis of 3-(4-Benzo[b]thiophene-2-carbonyl)-1-piperazinyl-1-benzo[b]thiophen-2-yl-1-propanones. <i>New Hetero Bis-Ligands with Potential 5-HT1A Serotonergic Activity. Heterocycles</i> , 2008, 75, 1913.	0.4	13
23	1-BENZOYL-2-(2-NITROPHENYL)-1H-BENZIMIDAZOLE DERIVATIVES: A NOVEL APPROACH TO THE DEVELOPMENT OF NEW HIV-1 REVERSE TRANSCRIPTASE INHIBITORS. <i>Journal of the Chilean Chemical Society</i> , 2007, 52, .	0.5	6
24	Studies on quinones. Part 41: Synthesis and cytotoxicity of isoquinoline-containing polycyclic quinones. <i>Bioorganic and Medicinal Chemistry</i> , 2006, 14, 5003-5011.	1.4	31
25	Solvent-Free Synthesis of 6-Arylbenzimidazo[1,2-c]quinazolines under Microwave-Irradiation. <i>Synthesis</i> , 2004, 2004, 436-440.	1.2	21
26	Studies on quinones. Part 35: Access to antiprotozoal active euryfurylquinones and hydroquinones. <i>Tetrahedron</i> , 2002, 58, 881-886.	1.0	28
27	Studies on quinones. Part 34: The reaction of styrene with activated 1,4-benzoquinones: access to potential antiprotozoal pyranobenzoquinones. <i>Tetrahedron</i> , 2001, 57, 8653-8658.	1.0	19
28	Regiospecific Michael reaction of (+)-euryfuran with activated 1,4-benzoquinones. <i>Tetrahedron Letters</i> , 2000, 41, 3563-3566.	0.7	6