

Michael J C Buckle

List of Publications by Year in descending order

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papers

695
citations

516215

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610482

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25
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1126
citing authors

#	ARTICLE	IF	CITATIONS
1	Analogues of 2- ² -hydroxychalcone with modified C4-substituents as the inhibitors against human acetylcholinesterase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021, 36, 130-137.	2.5	7
2	In vitro functional evaluation of isolaureline, dicentrine and glaucine enantiomers at 5-HT ₂ and 1 ₁ receptors. <i>Chemical Biology and Drug Design</i> , 2019, 93, 132-138.	1.5	12
3	2-Aryl-3-(arylideneamino)-1,2-dihydroquinazoline-4(3 <i>H</i>)-ones as inhibitors of cholinesterases and self-induced A β aggregation: biological evaluations and mechanistic insights from molecular dynamics simulations. <i>RSC Advances</i> , 2018, 8, 7818-7831.	1.7	6
4	Synthesis and evaluation of nuciferine and roemerine enantiomers as 5-HT ₂ and 1 ₁ receptor antagonists. <i>MedChemComm</i> , 2018, 9, 576-582.	3.5	12
5	Phosphodiesterase-5 inhibitors and their analogues as adulterants of herbal and food products: analysis of the Malaysian market, 2014-16. <i>Food Additives and Contaminants - Part A Chemistry, Analysis, Control, Exposure and Risk Assessment</i> , 2017, 34, 1101-1109.	1.1	24
6	Synthesis, Biological Evaluation and Molecular Modelling of 2-Hydroxychalcones as Acetylcholinesterase Inhibitors. <i>Molecules</i> , 2016, 21, 955.	1.7	24
7	In Silico and In Vitro Analysis of Bacoside A Aglycones and Its Derivatives as the Constituents Responsible for the Cognitive Effects of <i>Bacopa monnieri</i> . <i>PLoS ONE</i> , 2015, 10, e0126565.	1.1	60
8	Model studies on construction of the oxabicyclic [3.3.1] core of the mulberry Diels-Alder adducts morusalbanol A and 441772-64-1. <i>Tetrahedron Letters</i> , 2015, 56, 5082-5085.	0.7	5
9	Flavonoids with M1 Muscarinic Acetylcholine Receptor Binding Activity. <i>Molecules</i> , 2014, 19, 8933-8948.	1.7	19
10	Toward activated homology models of the human M1 muscarinic acetylcholine receptor. <i>Journal of Molecular Graphics and Modelling</i> , 2014, 49, 91-98.	1.3	13
11	Structure-Based Identification of Aporphines with Selective 5-HT _{2A} Receptor Binding Activity. <i>Chemical Biology and Drug Design</i> , 2013, 81, 250-256.	1.5	25
12	Induction of selective cytotoxicity and apoptosis in human T4-lymphoblastoid cell line (CEMss) by boesenbergin a isolated from <i>boesenbergia rotunda</i> rhizomes involves mitochondrial pathway, activation of caspase 3 and G2/M phase cell cycle arrest. <i>BMC Complementary and Alternative Medicine</i> , 2013, 13, 41.	3.7	35
13	Rational Discovery of Dengue Type 2 Non-Competitive Inhibitors. <i>Chemical Biology and Drug Design</i> , 2013, 82, 1-11.	1.5	38
14	Homology modeling of the human 5-HT _{1A} , 5-HT _{2A} , D ₁ , and D ₂ receptors: model refinement with molecular dynamics simulations and docking evaluation. <i>Journal of Molecular Modeling</i> , 2012, 18, 3639-3655.	0.8	26
15	Synthesis, Characterization, X-ray Crystallography, Acetyl Cholinesterase Inhibition and Antioxidant Activities of Some Novel Ketone Derivatives of Gallic Hydrazide-Derived Schiff Bases. <i>Molecules</i> , 2012, 17, 2408-2427.	1.7	30
16	Synthesis, Characterization, Acetylcholinesterase Inhibition, Molecular Modeling and Antioxidant Activities of Some Novel Schiff Bases Derived from 1-(2-Ketoiminoethyl)piperazines. <i>Molecules</i> , 2011, 16, 9316-9330.	1.7	16
17	Synthesis of (\pm)-kuwanon V and (\pm)-dorsterone methyl ethers via Diels-Alder reaction. <i>Tetrahedron Letters</i> , 2011, 52, 1797-1799.	0.7	32
18	An efficient one-pot synthesis of flavones. <i>Tetrahedron Letters</i> , 2011, 52, 3120-3123.	0.7	30

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19	An efficient synthesis of (±)-panduratin A and (±)-isopanduratin A, inhibitors of dengue-2 viral activity. Tetrahedron Letters, 2010, 51, 495-498.	0.7	36
20	Antimicrobial screening of plants used for traditional medicine in the state of Perak, Peninsular Malaysia. FĀ-toterapĀ-Āç, 2004, 75, 68-73.	1.1	118
21	Accurate determinations of the extent to which the SE2? reactions of allyl-, allenyl- and propargylsilanes are stereospecifically anti. Organic and Biomolecular Chemistry, 2004, 2, 749.	1.5	48
22	Accurate determination of the extent to which the SE2â€² reactions of an allenylsilane are stereospecifically anti. Tetrahedron Letters, 1993, 34, 2383-2386.	0.7	40
23	Accurate determination of the extent to which an SE2â€² reaction of an allylsilane is anti. Tetrahedron Letters, 1992, 33, 4479-4482.	0.7	38