

Rungnapha Saeeng

List of Publications by Year in descending order

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46
papers

725
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times ranked

702
citing authors

#	ARTICLE	IF	CITATIONS
1	<i>In vitro</i> galactose-targeted study of RSPP050-loaded micelles against liver hepatocellular carcinoma. <i>Pharmaceutical Development and Technology</i> , 2022, 27, 379-388.	1.1	7
2	Discovery of C-12 dithiocarbamate andrographolide analogues as inhibitors of SARS-CoV-2 main protease: <i>In vitro</i> and <i>in silico</i> studies. <i>Computational and Structural Biotechnology Journal</i> , 2022, 20, 2784-2797.	1.9	4
3	siRNA Targeting Mcl-1 Potentiates the Anticancer Activity of Andrographolide Nanosuspensions via Apoptosis in Breast Cancer Cells. <i>Pharmaceutics</i> , 2022, 14, 1196.	2.0	4
4	New 1,2,3-Triazole-genipin Analogues and Their Anti-Alzheimer's Activity. <i>ACS Omega</i> , 2022, 7, 24302-24316.	1.6	11
5	Synthesis and cytotoxic activity of new 7-acetoxy-12-amino-14-deoxy andrographolide analogues. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 33, 127741.	1.0	3
6	A novel synthetic acanthoic acid analogues and their cytotoxic activity in cholangiocarcinoma cells. <i>Bioorganic and Medicinal Chemistry</i> , 2021, 29, 115886.	1.4	7
7	Hypervalent Iodine(III)-Mediated Tandem Oxidative Dearomatization/Aziridination of Phenolic Amines: Synthesis of Functionalized Unactivated Aziridines. <i>Chemistry - A European Journal</i> , 2021, 27, 8473-8478.	1.7	3
8	Design, Synthesis, Evaluation and Molecular Docking Studies of 1,6-Bis-Triazole-Linked Galactoside Derivatives as Potential Anticancer Agents. <i>ChemistrySelect</i> , 2021, 6, 8052-8057.	0.7	2
9	Synthesis of propargylamine mycophenolate analogues and their selective cytotoxic activity towards neuroblastoma SH-SY5Y cell line. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 45, 128135.	1.0	8
10	Synthesis and biological evaluation of 1,6-bis-triazole-2,3,4-tri-O-benzyl- β -D-glucopyranosides as a novel β -glucosidase inhibitor in the treatment of Type 2 diabetes. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 50, 128331.	1.0	8
11	Antitumor effect of acanthoic acid against primary effusion lymphoma via inhibition of FLIP. <i>Phytotherapy Research</i> , 2021, 35, 7018-7026.	2.8	0
12	Design, Synthesis and Evaluations of New 10-Triazolyl-1-methoxygenipin Analogues for Their Cytotoxicity to Cancer Cells. <i>ChemistrySelect</i> , 2020, 5, 9540-9546.	0.7	8
13	Design and synthesis of C-12 dithiocarbamate andrographolide analogues as an anticancer agent. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127263.	1.0	11
14	Acetylcholinesterase Inhibitor From <i>Tabernaemontana pandacaqui</i> Flowers. <i>Natural Product Communications</i> , 2020, 15, 1934578X2091148.	0.2	2
15	New class of alkynyl glycoside analogues as tyrosinase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127276.	1.0	4
16	One-pot synthesis of substituted-amino triazole-glycosides. <i>Carbohydrate Research</i> , 2019, 484, 107780.	1.1	1
17	Fabrication and characterization of andrographolide analogue (3A.1) nanosuspensions stabilized by amphiphilic chitosan derivatives for colorectal cancer therapy. <i>Journal of Drug Delivery Science and Technology</i> , 2019, 54, 101287.	1.4	16
18	Folate-Functionalized Amphiphilic Chitosan Polymeric Micelles Containing Andrographolide Analogue (3A.1) for Colorectal Cancer. <i>Key Engineering Materials</i> , 2019, 819, 15-20.	0.4	5

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19	Metal-free selective synthesis of 2-substituted benzimidazoles catalyzed by Brønsted acidic ionic liquid: Convenient access to one-pot synthesis of N-alkylated 1,2-disubstituted benzimidazoles. <i>Tetrahedron</i> , 2019, 75, 3543-3552.	1.0	36
20	Synthetic analogues of durantoside I from <i>Citharexylum spinosum</i> L. and their cytotoxic activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 1558-1561.	1.0	3
21	A silyl andrographolide analogue suppresses Wnt/ β -catenin signaling pathway in colon cancer. <i>Biomedicine and Pharmacotherapy</i> , 2018, 101, 414-421.	2.5	21
22	One-Pot Approach for the Synthesis of Bis-indole-1,4-disubstituted-1,2,3-triazoles. <i>Journal of Organic Chemistry</i> , 2018, 83, 13233-13242.	1.7	12
23	The anti-cancer activity of an andrographolide analogue functions through a GSK-3 β -independent Wnt/ β -catenin signaling pathway in colorectal cancer cells. <i>Scientific Reports</i> , 2018, 8, 7924.	1.6	24
24	Apoptosis Induction and Antimigratory Activity of Andrographolide Analog (3A.1)-Incorporated Self-Assembled Nanoparticles in Cancer Cells. <i>AAPS PharmSciTech</i> , 2018, 19, 3123-3133.	1.5	15
25	Anticancer Activity of A Silyl Andrographolide Analogue Mediated Through Wnt/ β -Catenin Signaling In Colon Cancer Cells. <i>FASEB Journal</i> , 2018, 32, 1b680.	0.2	0
26	Synthesis and cytotoxic activity of 14-deoxy-12-hydroxyandrographolide analogs. <i>Medicinal Chemistry Research</i> , 2017, 26, 1653-1663.	1.1	7
27	One-pot three steps cascade synthesis of novel isoandrographolide analogues and their cytotoxic activity. <i>European Journal of Medicinal Chemistry</i> , 2017, 138, 952-963.	2.6	12
28	Synthesis of 14-deoxy-11,12-didehydroandrographolide analogues as potential cytotoxic agents for cholangiocarcinoma. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 5139-5143.	1.0	14
29	Preparation and Characterizations of RSPPO50-Loaded Polymeric Micelles Using Poly(ethylene Terephthalate) (PET) as a Matrix. <i>Journal of Applied Polymer Science</i> , 2017, 139, 45000.	0.6	24
30	Inhibition of Topoisomerase III α and Induction of Apoptosis in Gastric Cancer Cells by 19-Triisopropyl Andrographolide. <i>Asian Pacific Journal of Cancer Prevention</i> , 2017, 18, 2845-2851.	0.5	7
31	Novel Synthetic Mono-triazole Glycosides Induce G0/G1 Cell-cycle Arrest and Apoptosis in Cholangiocarcinoma Cells. <i>Anticancer Research</i> , 2016, 36, 5965-5974.	0.5	4
32	Solubility enhancement and in vitro evaluation of PEG-b-PLA micelles as nanocarrier of semi-synthetic andrographolide analogue for cholangiocarcinoma chemotherapy. <i>Pharmaceutical Development and Technology</i> , 2015, 21, 1-8.	1.1	22
33	Induction of apoptosis in cholangiocarcinoma by an andrographolide analogue is mediated through topoisomerase II α inhibition. <i>European Journal of Pharmacology</i> , 2014, 723, 148-155.	1.7	29
34	Green synthesis and anti-inflammatory studies of a series of 1,1-bis(heteroaryl)alkane derivatives. <i>European Journal of Medicinal Chemistry</i> , 2014, 83, 561-568.	2.6	33
35	Inhibition of topoisomerase II α activity and induction of apoptosis in mammalian cells by semi-synthetic andrographolide analogues. <i>Investigational New Drugs</i> , 2013, 31, 320-332.	1.2	25
36	One-pot synthesis of O-glycosyl triazoles by O-glycosylation "click" reaction. <i>Carbohydrate Research</i> , 2013, 375, 79-89.	1.1	9

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37	12-Amino-andrographolide analogues: synthesis and cytotoxic activity. Archives of Pharmacal Research, 2013, 36, 1454-1464.	2.7	19
38	New substituted C-19-andrographolide analogues with potent cytotoxic activities. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 49-52.	1.0	59
39	An efficient method for the selective synthesis of 2-deoxy-2-iodo-glycosides by O-glycosidation of d-glucal using I ₂ /Cu(OAc) ₂ . Carbohydrate Research, 2010, 345, 2401-2407.	1.1	18
40	Stereoselective C-Alkynylation, Allenylation, and Prop-2-ynylation Leading to Sugar Glycosides. Chemistry Letters, 2006, 35, 552-557.	0.7	27
41	Synthesis of Silyllallene Glycosides and Diene-Diglycosides by C-Glycosidation of d-Glucal with 1,4-Bis(trimethylsilyl)-2-butyne. Organic Letters, 2005, 7, 1585-1588.	0.7	57
42	Iodine catalyzes C-glycosidation of d-glucal with silylacetylene. Tetrahedron Letters, 2003, 44, 6211-6215.	2.4	31
43	Different C-Glycosidation Products of Glucal with Alkynyl or Propargyl Silanes under Acidic Conditions. Organic Letters, 2003, 5, 4883-4885.	0.4	22
44	Synthesis of Ciguatoxin (2S,5R)-ABC Segment. Heterocycles, 2001, 54, 789.	0.7	55
45	Partial synthesis of ciguatoxin (5R)-ABC segment. Tetrahedron Letters, 1999, 40, 1911-1914.	0.7	15
46	Electronic Factors in the C-Glycosidation with Silylacetylene. Chemistry Letters, 1999, 28, 467-468.		