Sadanandan E Velu

List of Publications by Year in descending order

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63 papers 1,539 citations

257357 24 h-index 35 g-index

66 all docs 66
docs citations

66 times ranked 2025 citing authors

#	Article	IF	Citations
1	Oxidation of $\hat{l}\pm,\hat{l}^2$ -enones and alkenes with oxone and sodium halides: A convenient laboratory preparation of chlorine and bromine. Tetrahedron Letters, 1996, 37, 2377-2380.	0.7	116
2	Attenuation of Nonsense-Mediated mRNA Decay Enhances In Vivo Nonsense Suppression. PLoS ONE, 2013, 8, e60478.	1.1	89
3	<i>In vitro</i> and <iin i="" vivo<=""> Anticancer Activity of Novel Synthetic Makaluvamine Analogues. Clinical Cancer Research, 2009, 15, 3511-3518.</iin>	3.2	62
4	Biochemical and Epigenetic Insights into L-2-Hydroxyglutarate, a Potential Therapeutic Target in Renal Cancer. Clinical Cancer Research, 2018, 24, 6433-6446.	3.2	54
5	Analogs of the marine alkaloid makaluvamines: Synthesis, topoisomerase II inhibition, and anticancer activity. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 2890-2893.	1.0	51
6	Identification of novel inhibitors of bacterial surface enzyme Staphylococcus aureus Sortase A. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 380-385.	1.0	45
7	Honokiol inhibits the growth of head and neck squamous cell carcinoma by targeting epidermal growth factor receptor. Oncotarget, 2015, 6, 21268-21282.	0.8	43
8	Discovery and Characterization of Dual Inhibitors of MDM2 and NFAT1 for Pancreatic Cancer Therapy. Cancer Research, 2018, 78, 5656-5667.	0.4	42
9	Discovery and evaluation of nNav1.5 sodium channel blockers with potent cell invasion inhibitory activity in breast cancer cells. Bioorganic and Medicinal Chemistry, 2018, 26, 2428-2436.	1.4	40
10	Cyanobacterial Metabolite Calothrixins: Recent Advances in Synthesis and Biological Evaluation. Marine Drugs, 2016, 14, 17.	2.2	39
11	New small-molecule inhibitors of dihydrofolate reductase inhibit Streptococcus mutans. International Journal of Antimicrobial Agents, 2015, 46, 174-182.	1.1	38
12	Structureâ€based approach to pharmacophore identification, <i>in silico</i> screening, and threeâ€dimensional quantitative structure–activity relationship studies for inhibitors of <i>Trypanosoma cruzi</i> dihydrofolate reductase function. Proteins: Structure, Function and Bioinformatics, 2008, 73, 889-901.	1.5	37
13	A novel synthetic iminoquinone, BA-TPQ, as an anti-breast cancer agent: in vitro and in vivo activity and mechanisms of action. Breast Cancer Research and Treatment, 2010, 123, 321-331.	1.1	37
14	Highly efficient delivery of potent anticancer iminoquinone derivative by multilayer hydrogel cubes. Acta Biomaterialia, 2017, 58, 386-398.	4.1	37
15	Conjugate Addition Reactions of α-Aminoalkylcuprates with α,β-Alkenyl-, α,β-Alkynyl-, α,βâ^'β,γ-Allenyl-, and α,βâ^'γ,Î'-Dienyl Carboxylic Acid Derivatives, Nitriles, and Sulfoxides. Journal of Organic Chemistry, 2000, 65, 8715-8724.	1.7	36
16	Experimental Therapy of Ovarian Cancer with Synthetic Makaluvamine Analog: In Vitro and In Vivo Anticancer Activity and Molecular Mechanisms of Action. PLoS ONE, 2011, 6, e20729.	1.1	36
17	MDM2-NFAT1 dual inhibitor, MA242: Effective against hepatocellular carcinoma, independent of p53. Cancer Letters, 2019, 459, 156-167.	3.2	36
18	Synthesis and characterization of potent inhibitors of Trypanosoma cruzi dihydrofolate reductase. Bioorganic and Medicinal Chemistry, 2010, 18, 4056-4066.	1.4	34

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19	Total synthesis of calothrixins A and B via oxidative radical reaction of cyclohexenone with aminophenanthridinedione. Tetrahedron, 2014, 70, 5928-5933.	1.0	32
20	Inhibition of <i>Streptococcus mutans</i> Biofilms by the Natural Stilbene Piceatannol Through the Inhibition of Glucosyltransferases. ACS Omega, 2018, 3, 8378-8385.	1.6	31
21	Synthesis and antiproliferative activity of benzyl and phenethyl analogs of makaluvamines. Bioorganic and Medicinal Chemistry, 2008, 16, 2541-2549.	1.4	30
22	Structure-Based Discovery of Small Molecule Inhibitors of Cariogenic Virulence. Scientific Reports, 2017, 7, 5974.	1.6	29
23	Pyrrolidinobenzoic acid inhibitors of influenza virus neuraminidase: modifications of essential pyrrolidinone ring substituents. Bioorganic and Medicinal Chemistry, 2003, 11, 2739-2749.	1.4	28
24	FBA-TPQ, a novel marine-derived compound as experimental therapy for prostate cancer. Investigational New Drugs, 2010, 28, 234-241.	1.2	28
25	Synthesis and In Vitro Anti-Lung Cancer Activity of Novel 1, 3, 4, 8- Tetrahydropyrrolo [4, 3, 2-de]quinolin-8(1H)-o ne Alkaloid Analogs. Medicinal Chemistry, 2009, 5, 227-236.	0.7	25
26	(α-Aminoalkyl)cuprates Prepared from Soluble Copper(I) Salts: Conjugate Additions to \hat{l} ±, \hat{l} 2-Unsaturated Carboxylic Acid Derivatives. Journal of Organic Chemistry, 1997, 62, 3798-3799.	1.7	24
27	Tethered Dimers as NAD Synthetase Inhibitors with Antibacterial Activity. Journal of Medicinal Chemistry, 2003, 46, 3371-3381.	2.9	24
28	Hydroxychalcone inhibitors of Streptococcus mutans glucosyl transferases and biofilms as potential anticaries agents. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 3508-3513.	1.0	24
29	Mini-Review: Bmx Kinase Inhibitors for Cancer Therapy. Recent Patents on Anti-Cancer Drug Discovery, 2013, 8, 228-238.	0.8	24
30	Synthesis and structure activity relationship studies of novel Staphylococcus aureus Sortase A inhibitors. European Journal of Medicinal Chemistry, 2010, 45, 3752-3761.	2.6	23
31	Pharmacological and nutritional targeting of voltage-gated sodium channels in the treatment of cancers. IScience, 2021, 24, 102270.	1.9	23
32	Preclinical Evaluation of Anticancer Efficacy and Pharmacological Properties of FBA-TPQ, a Novel Synthetic Makaluvamine Analog. Marine Drugs, 2012, 10, 1138-1155.	2.2	21
33	Preclinical Pharmacology of BA-TPQ, a Novel Synthetic Iminoquinone Anticancer Agent. Marine Drugs, 2010, 8, 2129-2141.	2.2	20
34	Elevated levels of NO are localized to distal airways in asthma. Free Radical Biology and Medicine, 2011, 50, 1679-1688.	1.3	20
35	Antibacterial Nicotinamide Adenine Dinucleotide Synthetase Inhibitors:Â Amide- and Ether-Linked Tethered Dimers with α-Amino Acid End Groups. Journal of Medicinal Chemistry, 2007, 50, 2612-2621.	2.9	16
36	Channel Gating Regulation by the Cystic Fibrosis Transmembrane Conductance Regulator (CFTR) First Cytosolic Loop. Journal of Biological Chemistry, 2016, 291, 1854-1865.	1.6	16

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37	SF3B1 homeostasis is critical for survival and therapeutic response in T cell leukemia. Science Advances, 2022, 8, eabj8357.	4.7	16
38	Total synthesis of zyzzyanones A–D. Tetrahedron, 2013, 69, 4105-4113.	1.0	15
39	Mitochondrial thiol modification by a targeted electrophile inhibits metabolism in breast adenocarcinoma cells by inhibiting enzyme activity and protein levels. Redox Biology, 2016, 8, 136-148.	3.9	15
40	Discovery and development of synthetic tricyclic pyrroloquinone (TPQ) alkaloid analogs for human cancer therapy. Frontiers of Chemical Science and Engineering, 2016, 10, 1-15.	2.3	15
41	Tethered Dimer Inhibitors of NAD Synthetase:Â Parallel Synthesis of an Aryl-Substituted SAR Library. ACS Combinatorial Science, 2005, 7, 898-904.	3.3	14
42	A facile synthesis of bispyrroloquinone and bispyrroloiminoquinone ring system of marine alkaloids. Tetrahedron Letters, 2009, 50, 3074-3076.	0.7	14
43	Regioselective Control in the Reactions of α-Aminoalkylcuprates with Allylic Substrates. Synlett, 1997, 1997, 1114-1116.	1.0	12
44	Recent advances in isolation, synthesis, and evaluation of bioactivities of bispyrroloquinone alkaloids of marine origin. Chinese Journal of Natural Medicines, 2015, 13, 561-577.	0.7	12
45	Nonsense-mediated decay controls the reactivation of the oncogenic herpesviruses EBV and KSHV. PLoS Biology, 2021, 19, e3001097.	2.6	12
46	Antibacterial and Antibiofilm Activities of Makaluvamine Analogs. Microorganisms, 2014, 2, 128-139.	1.6	11
47	A Novel Class of Mitochondria-Targeted Soft Electrophiles Modifies Mitochondrial Proteins and Inhibits Mitochondrial Metabolism in Breast Cancer Cells through Redox Mechanisms. PLoS ONE, 2015, 10, e0120460.	1.1	11
48	Toll-like receptor 9 expression is associated with breast cancer sensitivity to the growth inhibitory effects of bisphosphonates <i>in vitro</i> and <i>in vivo</i> . Oncotarget, 2016, 7, 87373-87389.	0.8	11
49	Discovery of Potent Inhibitors of Streptococcus mutans Biofilm with Antivirulence Activity. ACS Medicinal Chemistry Letters, 2021, 12, 48-55.	1.3	10
50	A Novel Marine Natural Product Derived Pyrroloiminoquinone with Potent Activity against Skin Cancer Cells. Marine Drugs, 2019, 17, 443.	2.2	9
51	Identification of the ZAK-MKK4-JNK-TGFβ Signaling Pathway as a Molecular Target for Novel Synthetic Iminoquinone Anticancer Compound BA-TPQ. Current Cancer Drug Targets, 2013, 13, 651-660.	0.8	8
52	Total Synthesis of Secobatzelline B. Synthetic Communications, 2007, 37, 2399-2409.	1.1	7
53	Development and validation of an HPLC method for quantitation of BAâ€TPQ, a novel iminoquinone anticancer agent, and an initial pharmacokinetic study in mice. Biomedical Chromatography, 2011, 25, 628-634.	0.8	7
54	Prognostic and therapeutic value of the Hippo pathway, RABL6A, and p53-MDM2 axes in sarcomas. Oncotarget, 2021, 12, 740-755.	0.8	7

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55	Synthesis of Pyrroloquinones via a CAN Mediated Oxidative Free Radical Reaction of 1,3-Dicarbonyl Compounds with Aminoquinones. Journal of Chemistry, 2013, 2013, 1-12.	0.9	6
56	Identification of TrkB Binders from Complex Matrices Using a Magnetic Drug Screening Nanoplatform. ACS Applied Bio Materials, 2021, 4, 6244-6255.	2.3	5
57	Azide-Mediated Detosylation of N-Tosylpyrroloiminoquinones and N-Tosylindole-4,7-quinones. Synlett, 2008, 2008, 2864-2868.	1.0	3
58	Synthetic Makaluvamine Analogs Decrease c-Kit Expression and Are Cytotoxic to Neuroendocrine Tumor Cells. Molecules, 2020, 25, 4940.	1.7	3
59	Synthesis of E Isomer and Crystal Structures of E and Z Isomers of 3-(2,5-Dimethoxyphenyl)-2-(4-methoxyphenyl)acrylonitrile. Journal of Chemical Crystallography, 2008, 38, 205-209.	0.5	2
60	Development and validation of an HPLC-MS/MS analytical method for quantitative analysis of TCBA-TPQ, a novel anticancer makaluvamine analog, and application in a pharmacokinetic study in rats. Chinese Journal of Natural Medicines, 2015, 13, 554-560.	0.7	2
61	Synthesis, Separation and Crystal Structures of E and Z Isomers of 3-(2,5-Dimethoxyphenyl)-2-(4-Methoxyphenyl)Acrylic Acid. Journal of Chemical Crystallography, 2008, 38, 189-194.	0.5	1
62	An expedient synthesis of murrayaquinone A via a novel oxidative free radical reaction. Tetrahedron Letters, 2018, 59, 550-553.	0.7	1
63	Synthesis and the Crystal Structure of (E)-2-(7-(3-(Thiophen-2-yl)acrylamido)-2,3-dihydro-5-oxobenzo[e][1,4]oxazepin-1(5H)-yl)ethyl acetate. Journal of Chemical Crystallography, 2009, 39, 902.	0.5	0