Pradeep B Lukka

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1453798/publications.pdf

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		1464605	1526636	
17	146	7	10	
papers	citations	h-index	g-index	
18	18	18	167	
all docs	docs citations	times ranked	citing authors	

#	Article	IF	CITATIONS
1	Colchicine-Binding Site Agent CH-2-77 as a Potent Tubulin Inhibitor Suppressing Triple-Negative Breast Cancer. Molecular Cancer Therapeutics, 2022, 21, 1103-1114.	1.9	5
2	Discovery of <i>N</i> -(3,4-Dimethylphenyl)-4-(4-isobutyrylphenyl)-2,3,3a,4,5,9b-hexahydrofuro[3,2- <i>c</i>)quinoline-8-sulfor as a Potent Dual MDM2/XIAP Inhibitor. Journal of Medicinal Chemistry, 2021, 64, 1930-1950.	na zni de	10
3	Use of Realâ€World Data and Pharmacometric Modeling in Support of Lacosamide Dosing in Pediatric Patients Under 4 Years of Age. Journal of Clinical Pharmacology, 2021, 61, 881-888.	1.0	9
4	Model-Based Exposure-Response Assessment for Spectinamide 1810 in a Mouse Model of Tuberculosis. Antimicrobial Agents and Chemotherapy, 2021, 65, e0174420.	1.4	7
5	X-ray Crystallography-Guided Design, Antitumor Efficacy, and QSAR Analysis of Metabolically Stable Cyclopenta-Pyrimidinyl Dihydroquinoxalinone as a Potent Tubulin Polymerization Inhibitor. Journal of Medicinal Chemistry, 2021, 64, 13072-13095.	2.9	13
6	Preclinical Evaluation of Inhalational Spectinamide-1599 Therapy against Tuberculosis. ACS Infectious Diseases, 2021, 7, 2850-2863.	1.8	8
7	Development and Characterization of a Dry Powder Formulation for Anti-Tuberculosis Drug Spectinamide 1599. Pharmaceutical Research, 2019, 36, 136.	1.7	19
8	Comparative pharmacokinetics of spectinamide 1599 after subcutaneous and intrapulmonary aerosol administration in mice. Tuberculosis, 2019, 114, 119-122.	0.8	8
9	Sterilization of Mycobacterium tuberculosis infected samples using methanol preserves anti-tuberculosis drugs for subsequent pharmacological testing studies. Tuberculosis, 2019, 117, 52-55.	0.8	2
10	Primary Lung Dendritic Cell Cultures to Assess Efficacy of Spectinamide-1599 Against Intracellular Mycobacterium tuberculosis. Frontiers in Microbiology, 2018, 9, 1895.	1.5	5
11	Alterations in cellular pharmacokinetics and pharmacodynamics of elvitegravir in response to ethanol exposure in HIV-1 infected monocytic (U1) cells. PLoS ONE, 2017, 12, e0172628.	1.1	15
12	Selective cellular uptake and retention of SN 28049, a new DNA-binding topoisomerase II-directed antitumor agent. Cancer Chemotherapy and Pharmacology, 2014, 74, 25-35.	1.1	8
13	Tumour tissue selectivity in the uptake and retention of SN 28049, a new topoisomerase II-directed anticancer agent. Cancer Chemotherapy and Pharmacology, 2013, 72, 1013-1022.	1.1	4
14	Comparison of a homologous series of benzonaphthyridine anti-cancer agents in mice: divergence between tumour and plasma pharmacokinetics. Cancer Chemotherapy and Pharmacology, 2012, 70, 151-160.	1.1	8
15	A rapid LC–MS/MS method for the quantitation of a series of benzonaphthyridine derivatives: Application to in vivo pharmacokinetic and lipophilicity studies in drug development. Journal of Pharmaceutical and Biomedical Analysis, 2012, 63, 9-16.	1.4	7
16	Pharmacokinetics and distribution of SN 28049, a novel DNA binding anticancer agent, in mice. Cancer Chemotherapy and Pharmacology, 2010, 65, 1145-1152.	1.1	10
17	Development and validation of a liquid chromatographya—mass spectrometry (LCa—MS) assay for the determination of the anti-cancer agent N-[2-(dimethylamino)ethyl]-2,6-dimethyl-1-oxo-1,2-dihydrobenzo[b]-1,6-naphthyridine-4-carboxamide (SN) Tj ETC	<u>)</u> q 1. 1 0.78	34 3 14 rgBT /(