## Sonali S Bharate

List of Publications by Year in descending order

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SONALLS RHARATE

#	Article	IF	CITATIONS
1	Non-enzymatic browning in citrus juice: chemical markers, their detection and ways to improve product quality. Journal of Food Science and Technology, 2014, 51, 2271-2288.	1.4	101
2	Modulation of Thermoreceptor TRPM8 by Cooling Compounds. ACS Chemical Neuroscience, 2012, 3, 248-267.	1.7	81
3	Critical Analysis of Drug Product Recalls due to Nitrosamine Impurities. Journal of Medicinal Chemistry, 2021, 64, 2923-2936.	2.9	78
4	Why Are the Majority of Active Compounds in the CNS Domain Natural Products? A Critical Analysis. Journal of Medicinal Chemistry, 2018, 61, 10345-10374.	2.9	67
5	Discovery of a marine-derived bis-indole alkaloid fascaplysin, as a new class of potent P-glycoprotein inducer and establishment of its structure–activity relationship. European Journal of Medicinal Chemistry, 2016, 107, 1-11.	2.6	66
6	<i>Crocus sativus</i> Extract Tightens the Blood-Brain Barrier, Reduces Amyloid β Load and Related Toxicity in 5XFAD Mice. ACS Chemical Neuroscience, 2017, 8, 1756-1766.	1.7	66
7	Crocetin promotes clearance of amyloid-β by inducing autophagy via the STK11/LKB1-mediated AMPK pathway. Autophagy, 2021, 17, 3813-3832.	4.3	62
8	Recent developments in pharmaceutical salts: FDA approvals from 2015 to 2019. Drug Discovery Today, 2021, 26, 384-398.	3.2	55
9	Determining Partition Coefficient (Log P), Distribution Coefficient (Log D) and Ionization Constant (pKa) in Early Drug Discovery. Combinatorial Chemistry and High Throughput Screening, 2016, 19, 461-469.	0.6	48
10	Discovery and characterization of novel CYP1B1 inhibitors based on heterocyclic chalcones: Overcoming cisplatin resistance in CYP1B1-overexpressing lines. European Journal of Medicinal Chemistry, 2017, 129, 159-174.	2.6	41
11	Pyrano-isochromanones as IL-6 Inhibitors: Synthesis, in Vitro and in Vivo Antiarthritic Activity. Journal of Medicinal Chemistry, 2014, 57, 7085-7097.	2.9	39
12	6-Aryl substituted 4-(4-cyanomethyl) phenylamino quinazolines as a new class of isoform-selective PI3K-alpha inhibitors. European Journal of Medicinal Chemistry, 2016, 122, 731-743.	2.6	39
13	Design of Novel 3-Pyrimidinylazaindole CDK2/9 Inhibitors with Potent In Vitro and In Vivo Antitumor Efficacy in a Triple-Negative Breast Cancer Model. Journal of Medicinal Chemistry, 2017, 60, 9470-9489.	2.9	39
14	Discovery and Preclinical Development of IIIM-290, an Orally Active Potent Cyclin-Dependent Kinase Inhibitor. Journal of Medicinal Chemistry, 2018, 61, 1664-1687.	2.9	39
15	Analysis of clinical trials on biomaterial and therapeutic applications of chitosan: A review. Carbohydrate Polymers, 2022, 278, 118999.	5.1	39
16	Impact of preformulation on drug development. Expert Opinion on Drug Delivery, 2013, 10, 1239-1257.	2.4	37
17	Thermodynamic equilibrium solubility measurements in simulated fluids by 96-well plate method in early drug discovery. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 1561-1567.	1.0	34
18	Nitrofuranyl Methyl Piperazines as New Anti-TB Agents: Identification, Validation, Medicinal Chemistry, and PK Studies. ACS Medicinal Chemistry Letters, 2015, 6, 1041-1046.	1.3	33

SONALI S BHARATE

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19	Biphenyl-4-carboxylic Acid [2-(1 <i>H</i> -Indol-3-yl)-ethyl]-methylamide (CA224), a Nonplanar Analogue of Fascaplysin, Inhibits Cdk4 and Tubulin Polymerization: Evaluation of in Vitro and in Vivo Anticancer Activity. Journal of Medicinal Chemistry, 2014, 57, 9658-9672.	2.9	32
20	Anti-inflammatory chromone alkaloids and glycoside from Dysoxylum binectariferum. Tetrahedron Letters, 2017, 58, 3974-3978.	0.7	32
21	A chromatography-free isolation of rohitukine from leaves of Dysoxylum binectariferum: Evaluation for in vitro cytotoxicity, Cdk inhibition and physicochemical properties. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 3457-3463.	1.0	31
22	Colchicine derivatives with potent anticancer activity and reduced P-glycoprotein induction liability. Organic and Biomolecular Chemistry, 2015, 13, 5674-5689.	1.5	30
23	Synthesis and anti-proliferative activities of new derivatives of embelin. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 4865-4870.	1.0	28
24	Preclinical Development of Crocus sativus-Based Botanical Lead IIIM-141 for Alzheimer's Disease: Chemical Standardization, Efficacy, Formulation Development, Pharmacokinetics, and Safety Pharmacology. ACS Omega, 2018, 3, 9572-9585.	1.6	26
25	Synthesis, pH dependent, plasma and enzymatic stability of bergenin prodrugs for potential use against rheumatoid arthritis. Bioorganic and Medicinal Chemistry, 2017, 25, 5513-5521.	1.4	25
26	Binary and ternary solid dispersions of an anticancer preclinical lead, IIIM-290: In vitro and in vivo studies. International Journal of Pharmaceutics, 2019, 570, 118683.	2.6	25
27	Chemical analysis of saffron by HPLC based crocetin estimation. Journal of Pharmaceutical and Biomedical Analysis, 2020, 181, 113094.	1.4	25
28	Discovery of 7-(Prolinol-N-yl)-2-phenylamino-thiazolo[5,4-d]pyrimidines as Novel Non-Nucleoside Partial Agonists for the A2A Adenosine Receptor: Prediction from Molecular Modeling. Journal of Medicinal Chemistry, 2016, 59, 5922-5928.	2.9	23
29	Modulating lipophilicity of rohitukine via prodrug approach: Preparation, characterization, and in vitro enzymatic hydrolysis in biorelevant media. European Journal of Pharmaceutical Sciences, 2016, 92, 203-211.	1.9	19
30	Carboxylic Acid Counterions in FDA-Approved Pharmaceutical Salts. Pharmaceutical Research, 2021, 38, 1307-1326.	1.7	17
31	Quantitative Determination and Characterization of a Kashmir Saffron ( <i>Crocus sativus</i> ) Tj ETQq1 1 0.784 and HPTLC Investigations. ACS Omega, 2021, 6, 23460-23474.	·314 rgBT 1.6	/Overlock 10 15
32	Synthesis and Biological Evaluation of Polar Functionalities Containing Nitrodihydroimidazooxazoles as Anti-TB Agents. ACS Medicinal Chemistry Letters, 2015, 6, 1059-1064.	1.3	12
33	Selection of a Water-Soluble Salt Form of a Preclinical Candidate, IIIM-290: Multiwell-Plate Salt Screening and Characterization. ACS Omega, 2018, 3, 8365-8377.	1.6	12
34	Modulation of biopharmaceutical properties of acidic drugs using cationic counterions: A critical analysis of FDA-approved pharmaceutical salts. International Journal of Pharmaceutics, 2021, 607, 120993.	2.6	12
35	Antidiabetic potential of polyherbal formulation DB14201: Preclinical development, safety and efficacy studies. Journal of Ethnopharmacology, 2017, 197, 218-230.	2.0	11
36	Orally Effective Aminoalkyl 10 H â€Indolo[3,2―b ]quinolineâ€11â€carboxamide Kills the Malaria Parasite by Inhibiting Host Hemoglobin Uptake. ChemMedChem, 2018, 13, 2581-2598.	1.6	11

SONALI S BHARATE

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37	Discovery and preclinical development of IIIM-160, a Bergenia ciliata-based anti-inflammatory and anti-arthritic botanical drug candidate. Journal of Integrative Medicine, 2019, 17, 192-204.	1.4	11
38	Recent developments in the management of Huntington's disease. Bioorganic Chemistry, 2022, 120, 105642.	2.0	11
39	Phosphate moiety in FDAâ€approved pharmaceutical salts and prodrugs. Drug Development Research, 2022, 83, 1059-1074.	1.4	10
40	Enhancing Biopharmaceutical Attributes of Khellin by Amorphous Binary Solid Dispersions. AAPS PharmSciTech, 2021, 22, 260.	1.5	8
41	Trifluoroacetic acid catalyzed thiophenylmethylation and thioalkylmethylation of lactams and phenols via domino three-component reaction in water. RSC Advances, 2014, 4, 14081-14088.	1.7	6
42	Design, synthesis and P-gp induction activity of aryl phosphonate esters: identification of tetraethyl-2-phenylethene-1,1-diyldiphosphonate as an orally bioavailable P-gp inducer. MedChemComm, 2016, 7, 1910-1915.	3.5	6
43	Impurity profiling of anticancer preclinical candidate, IIIM-290. Journal of Pharmaceutical and Biomedical Analysis, 2019, 166, 1-5.	1.4	6
44	Evaluation of rohitukine-enriched fraction of Dysoxylum binectariferum Hook.f. (leaves) as anti-arthritic phytopharmaceutical candidate: Chemical standardization, in-vivo validation, formulation development and oral pharmacokinetics. Journal of Ethnopharmacology, 2020, 254, 112758.	2.0	6
45	Identification of plant-based multitargeted leads for Alzheimer's disease: In-vitro and in-vivo validation of Woodfordia fruticosa (L.) Kurz. Phytomedicine, 2021, 91, 153659.	2.3	6
46	Engineering solid dispersions of anticancer preclinical lead, IIIM-985: Physicochemical characterization and in vivo pharmacokinetics. Journal of Drug Delivery Science and Technology, 2019, 49, 594-602.	1.4	5
47	Thermodynamic solubility determination of khellin in eight mono-solvents at the range of 298.15 to 323.15ÂK. Journal of Molecular Liquids, 2022, 351, 118637.	2.3	5
48	Modulation of biopharmaceutical properties of drugs using sulfonate counterions: A critical analysis of FDA-approved pharmaceutical salts. Journal of Drug Delivery Science and Technology, 2021, 66, 102913.	1.4	4
49	Analytical Methods for Furanochromone Natural Product, Khellin and Its Inspired Drug Candidates, Amiodarone and Sodium Cromoglycate. Critical Reviews in Analytical Chemistry, 0, , 1-16.	1.8	0