Jingshan Shen

List of Publications by Year in descending order

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687220 289141 1,789 47 13 40 citations h-index g-index papers 48 48 48 3543 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	"One-Pot―Synthesis of Molnupiravir from Cytidine. Organic Process Research and Development, 2022, 26, 358-364.	1.3	10
2	Synthesis and antiviral activity of 2′â€deoxyâ€6′â€substituted carbocyclic nucleosides. Chemical Biology ar Drug Design, 2022, 99, 561-572.	nd _{1.5}	1
3	Artemisinin derivative TPN10466 suppresses immune cell migration and Th1/Th17 differentiation to ameliorate disease severity in experimental autoimmune encephalomyelitis. Cellular Immunology, 2022, 373, 104500.	1.4	7
4	Oral remdesivir derivative VV116 is a potent inhibitor of respiratory syncytial virus with efficacy in mouse model. Signal Transduction and Targeted Therapy, 2022, 7, 123.	7.1	14
5	TPN171H alleviates pulmonary hypertension via inhibiting inflammation in hypoxia and monocrotaline-induced rats. Vascular Pharmacology, 2022, 145, 107017.	1.0	3
6	Synthesis and anti-SARS-CoV-2 activity of deuterated GS-441524 analogs. Tetrahedron Letters, 2022, 104, 154012.	0.7	6
7	Challenges and stepwise fit-for-purpose optimization for bioanalyses of remdesivir metabolites nucleotide monophosphate and triphosphate in mouse tissues using LC–MS/MS. Journal of Pharmaceutical and Biomedical Analysis, 2021, 194, 113806.	1.4	9
8	One step stereoselective synthesis of oxazoline-fused saccharides and their conversion into the corresponding 1,2-cis glycosylamines bearing various protected groups. Organic and Biomolecular Chemistry, 2021, 19, 1580-1588.	1.5	1
9	Chronic administration of synthetic cannabidiol induces antidepressant effects involving modulation of serotonin and noradrenaline levels in the hippocampus. Neuroscience Letters, 2021, 744, 135594.	1.0	15
10	Significant Inhibition of Porcine Epidemic Diarrhea Virus In Vitro by Remdesivir, Its Parent Nucleoside and \hat{l}^2 -d-N4-hydroxycytidine. Virologica Sinica, 2021, 36, 997-1005.	1.2	12
11	9,10-Anhydrodehydroartemisinin Attenuates Experimental Autoimmune Encephalomyelitis by Inhibiting Th1 and Th17 Cell Differentiation. Inflammation, 2021, 44, 1793-1802.	1.7	11
12	Weinreb Amide Approach to the Practical Synthesis of a Key Remdesivir Intermediate. Journal of Organic Chemistry, 2021, 86, 5065-5072.	1.7	15
13	Identification of pyrogallol as a warhead in design of covalent inhibitors for the SARS-CoV-2 3CL protease. Nature Communications, 2021, 12, 3623.	5.8	111
14	SARS-CoV-2 envelope protein causes acute respiratory distress syndrome (ARDS)-like pathological damages and constitutes an antiviral target. Cell Research, 2021, 31, 847-860.	5.7	102
15	A Phase I Study to Evaluate the Safety, Tolerability, and Pharmacokinetics of TPN171H, a Novel Phosphodiesterase Type 5 Inhibitor, in Healthy Subjects. Drug Design, Development and Therapy, 2021, Volume 15, 2947-2959.	2.0	3
16	Potency and pharmacokinetics of GS-441524 derivatives against SARS-CoV-2. Bioorganic and Medicinal Chemistry, 2021, 46, 116364.	1.4	21
17	Design and development of an oral remdesivir derivative VV116 against SARS-CoV-2. Cell Research, 2021, 31, 1212-1214.	5.7	71
18	Scalable Process for Making 5,7-Dichlorotetrahydroisoquinoline-6-carboxylic Acid Using Methylene as the Protecting Group. Organic Process Research and Development, 2021, 25, 2447-2452.	1.3	4

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19	Synthesis of CBD and Its Derivatives Bearing Various C4′-Side Chains with a Late-Stage Diversification Method. Journal of Organic Chemistry, 2020, 85, 2704-2715.	1.7	31
20	The novel small-molecule TPN10456 inhibits Th17 cell differentiation and protects against experimental autoimmune encephalomyelitis. Cellular and Molecular Immunology, 2020, 17, 1290-1293.	4.8	2
21	Targeted Drugs for Treatment of Pulmonary Arterial Hypertension: Past, Present, and Future Perspectives. Journal of Medicinal Chemistry, 2020, 63, 15153-15186.	2.9	20
22	Synthesis and biological investigation of triazolopyridinone derivatives as potential multireceptor atypical antipsychotics. Bioorganic and Medicinal Chemistry Letters, 2020, 30, 127027.	1.0	5
23	Structural basis for inhibition of the RNA-dependent RNA polymerase from SARS-CoV-2 by remdesivir. Science, 2020, 368, 1499-1504.	6.0	950
24	Improved Synthesis of 6-Chloro-5-methylpyridin-2-amine: A Key Intermediate for Making Lumacaftor. Organic Process Research and Development, 2020, 24, 1175-1179.	1.3	4
25	An improved synthesis of telmisartan <i>via</i> the copper-catalyzed cyclization of <i>o</i> -haloarylamidines. RSC Advances, 2020, 10, 13717-13721.	1.7	8
26	Nature brings new avenues to the therapy of central nervous system diseasesâ€"An overview of possible treatments derived from natural products. Science China Life Sciences, 2019, 62, 1332-1367.	2.3	20
27	Continuation of structure–activity relationship study of novel benzamide derivatives as potential antipsychotics. Archiv Der Pharmazie, 2019, 352, 1800306.	2.1	2
28	Oxidative Aromatization of 3,4-Dihydroquinolin- $2(1 < i > H < / i >)$ -ones to Quinolin- $2(1 < i > H < / i >)$ -ones Using Transition-Metal-Activated Persulfate Salts. Journal of Organic Chemistry, 2019, 84, 8702-8709.	1.7	10
29	Pharmacokinetics-Driven Optimization of 4(3 <i>H</i>)-Pyrimidinones as Phosphodiesterase Type 5 Inhibitors Leading to TPN171, a Clinical Candidate for the Treatment of Pulmonary Arterial Hypertension. Journal of Medicinal Chemistry, 2019, 62, 4979-4990.	2.9	25
30	Industry-Oriented Route Evaluation and Process Optimization for the Preparation of Brexpiprazole. Organic Process Research and Development, 2019, 23, 852-857.	1.3	5
31	Synthesis and Biological Evaluation of Fiveâ€Atomâ€Linkerâ€Based Arylpiperazine Derivatives with an Atypical Antipsychotic Profile. ChemMedChem, 2019, 14, 2042-2051.	1.6	6
32	Rational design of 5-((1H-imidazol-1-yl)methyl)quinolin-8-ol derivatives as novel bromodomain-containing protein 4 inhibitors. European Journal of Medicinal Chemistry, 2019, 163, 281-294.	2.6	13
33	Discovery of pyrimidine nucleoside dual prodrugs and pyrazine nucleosides as novel anti-HCV agents. Bioorganic and Medicinal Chemistry, 2019, 27, 748-759.	1.4	6
34	A Facile Epoxide Aminolysis Promoted by (<i>t</i> -BuO) ₂ Mg and Its Application to the Synthesis of Efinaconazole. Organic Process Research and Development, 2018, 22, 625-632.	1.3	5
35	Synthesis and biological evaluation of a series of novel pyridinecarboxamides as potential multi-receptor antipsychotic drugs. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 606-611.	1.0	8
36	Aminobenzisoxazole compounds as agonists of $\hat{l}\pm7$ nicotinic acetylcholine receptors: a patent evaluation (WOÂ2017027600). Expert Opinion on Therapeutic Patents, 2018, 28, 429-436.	2.4	0

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37	Synthesis and biological investigation of tetrahydropyridopyrimidinone derivatives as potential multireceptor atypical antipsychotics. Bioorganic and Medicinal Chemistry, 2017, 25, 4904-4916.	1.4	11
38	Synthesis, structure–activity relationships, and biological evaluation of a series of benzamides as potential multireceptor antipsychotics. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 3141-3147.	1.0	6
39	Facile Synthesis of Substituted 4-Alkoxy-2-oxazolines and Exploration of the Reaction Mechanism. Synthesis, 2016, 48, 1331-1343.	1.2	5
40	An Improved Synthesis of 4-(1-Piperazinyl)benzo[<i>b</i>]thiophene Dihydrochloride. Organic Process Research and Development, 2015, 19, 555-558.	1.3	15
41	Thermodynamic and Structural Characterization of Halogen Bonding in Protein–Ligand Interactions: A Case Study of PDE5 and Its Inhibitors. Journal of Medicinal Chemistry, 2014, 57, 3588-3593.	2.9	37
42	Exploration of the 5-bromopyrimidin-4(3H)-ones as potent inhibitors of PDE5. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 4944-4947.	1.0	9
43	Design, Synthesis, and Pharmacological Evaluation of Monocyclic Pyrimidinones as Novel Inhibitors of PDE5. Journal of Medicinal Chemistry, 2012, 55, 10540-10550.	2.9	28
44	Utilization of Halogen Bond in Lead Optimization: A Case Study of Rational Design of Potent Phosphodiesterase Type 5 (PDE5) Inhibitors. Journal of Medicinal Chemistry, 2011, 54, 5607-5611.	2.9	108
45	2-Phenylquinazolin-4(3H)-one, a class of potent PDE5 inhibitors with high selectivity versus PDE6. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 2777-2779.	1.0	20
46	An Improved Synthetic Route for Preparative Process of Vardenafil. Organic Process Research and Development, 2009, 13, 1206-1208.	1.3	7
47	Facile and Cost-Effective Route for the Synthesis of Simmerafil. Organic Process Research and Development, 0, , .	1.3	0