Douglas C Beshore

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2272798/publications.pdf

Version: 2024-02-01

20 papers 548

840776 11 h-index 18 g-index

22 all docs 22 docs citations

times ranked

22

786 citing authors

#	Article	IF	Citations
1	Recent advances in the synthesis of diketopiperazines. Tetrahedron, 2002, 58, 3297-3312.	1.9	190
2	Design and Synthesis of Novel Isoquinoline-3-nitriles as Orally Bioavailable Kv1.5 Antagonists for the Treatment of Atrial Fibrillation. Journal of Medicinal Chemistry, 2006, 49, 6954-6957.	6.4	95
3	SYNTHESES AND TRANSFORMATIONS OF PIPERAZINONE RINGS. A REVIEW. Organic Preparations and Procedures International, 2002, 34, 367-404.	1.3	44
4	Preparation of Substituted Piperazinones via Tandem Reductive Aminationâ^'(N,Nâ€~-Acyl) Tj ETQq0 0 0 rgBT /Ov	verlock 10 4.6) Tf 50 622 Td
5	Atrial Antifibrillatory Effects of Structurally Distinct IKur Blockers 3-[(Dimethylamino)methyl]-6-methoxy-2-methyl-4-phenylisoquinolin-1(2H)-one and 2-Phenyl-1,1-dipyridin-3-yl-2-pyrrolidin-1-yl-ethanol in Dogs with Underlying Heart Failure. Journal of Pharmacology and Experimental Therapeutics. 2008. 324. 322-330.	2.5	32
6	MK-7622: A First-in-Class M ₁ Positive Allosteric Modulator Development Candidate. ACS Medicinal Chemistry Letters, 2018, 9, 652-656.	2.8	25
7	Discovery of ethyl ketone-based HDACs 1, 2, and 3 selective inhibitors for HIV latency reactivation. Bioorganic and Medicinal Chemistry Letters, 2020, 30, 127197.	2.2	19
8	Informing the Selection of Screening Hit Series with in Silico Absorption, Distribution, Metabolism, Excretion, and Toxicity Profiles. Journal of Medicinal Chemistry, 2017, 60, 6771-6780.	6.4	17
9	Microbial biotransformation – an important tool for the study of drug metabolism. Xenobiotica, 2019, 49, 877-886.	1.1	17
10	Efficient synthesis of unsymmetrical 1,4-disubstituted-2,3-diketopiperazines via tandem reductive amination–cyclization. Tetrahedron Letters, 2000, 41, 8735-8739.	1.4	15
11	Discovery of triarylethanolamine inhibitors of the Kv1.5 potassium channel. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 2493-2496.	2.2	13
12	SAR Studies on Carboxylic Acid Series M ₁ Selective Positive Allosteric Modulators (PAMs). Current Topics in Medicinal Chemistry, 2014, 14, 1738-1754.	2.1	11
13	Preparation of 2,4,5-trisubstituted pyrazolo[4,3-c]quinolin-3-ones. Tetrahedron Letters, 2010, 51, 970-973.	1.4	10
14	Preparation of Ethyl 5-lodo-1H-indole- 2-carboxylate. Synthetic Communications, 2003, 33, 2423-2427.	2.1	9
15	Redefining the Histone Deacetylase Inhibitor Pharmacophore: High Potency with No Zinc Cofactor Interaction. ACS Medicinal Chemistry Letters, 2021, 12, 540-547.	2.8	9
16	Evaluation of amino acid-based linkers in potent macrocyclic inhibitors of farnesyl-protein transferase. Bioorganic and Medicinal Chemistry Letters, 2001, 11, 1817-1821.	2.2	6
17	2,5â€Disubstituted Pyrazolo[4,3â€ <i>c</i>]cinnolinâ€3â€ones. Journal of Heterocyclic Chemistry, 2017, 54, 1558-1571.	2.6	1
18	Building a Culture of Medicinal Chemistry Knowledge Sharing. Journal of Medicinal Chemistry, 2022, 65, 3776-3785.	6.4	1

#	Article	IF	CITATIONS
19	Preparation of Ethyl 5-lodo-1H-indole-2-carboxylate ChemInform, 2003, 34, no.	0.0	0

Preparation of Substituted Piperazinones via Tandem Reductive Amination-(N,N′-Acyl) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702 Td (700 m) Preparation of Substituted Piperazinones via Tandem Reductive Amination-(N,N′-Acyl) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702 Td (700 m) Preparation of Substituted Piperazinones via Tandem Reductive Amination-(N,N′-Acyl) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702 Td (700 m) Preparation of Substituted Piperazinones via Tandem Reductive Amination-(N,N′-Acyl) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702 Td (700 m) Preparation of Substituted Piperazinones via Tandem Reductive Amination-(N,N′-Acyl) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702 Td (700 m) Preparation of Substituted Piperazinones via Tandem Reductive Amination-(N,N′-Acyl) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702 Td (700 m) Preparation of Substituted Piperazinones via Tandem Reductive Amination-(N,N′-Acyl) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702 Td (700 m) Preparation of Substituted Piperazinones via Tandem Reductive Amination-(N,N′-Acyl) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702 Td (700 m) Preparation of Substituted Piperazinones via Tandem Reductive Amination-(N,N′-Acyl) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702 Td (700 m) Preparation of Substituted Piperazinones via Tandem Reductive Amination of Substitu