

Laurent Knerr

List of Publications by Year in descending order

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Version: 2024-02-01

17
papers

470
citations

759233

12
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839539

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all docs

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docs citations

18
times ranked

654
citing authors

#	ARTICLE	IF	CITATIONS
1	A structure-specific small molecule inhibits a miRNA-200 family member precursor and reverses a type 2 diabetes phenotype. <i>Cell Chemical Biology</i> , 2022, 29, 300-311.e10.	5.2	13
2	Glucagon Like Peptide 1 Receptor Agonists for Targeted Delivery of Antisense Oligonucleotides to Pancreatic Beta Cell. <i>Journal of the American Chemical Society</i> , 2021, 143, 3416-3429.	13.7	39
3	Melanocortin 1 Receptor Agonists Based on a Bivalent, Bicyclic Peptide Framework. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 9906-9915.	6.4	6
4	Target-Directed Approaches for Screening Small Molecules against RNA Targets. <i>SLAS Discovery</i> , 2020, 25, 869-894.	2.7	23
5	Synthesis of Diverse β -Fluoroalkoxyaryl Derivatives and Their Use for the Generation of Fluorinated Macrocycles. <i>Chemistry - A European Journal</i> , 2019, 25, 1184-1187.	3.3	4
6	Stereocontrolled Synthesis of Tetrafluoropentanol: Multivincinal Fluorinated Alkane Units for Drug Discovery. <i>Organic Letters</i> , 2019, 21, 7741-7745.	4.6	20
7	Late-Stage Functionalization of Histidine in Unprotected Peptides. <i>Angewandte Chemie</i> , 2019, 131, 19272-19278.	2.0	34
8	Late-Stage Functionalization of Histidine in Unprotected Peptides. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 19096-19102.	13.8	47
9	Iridium-Catalyzed Asymmetric Hydrogenation of α -Alkyl β -Aryl Furan-Containing Imines: an Efficient Route to Unnatural α -Alkyl Arylalanines and Related Derivatives.. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 578-584.	4.3	12
10	Fluorine-Directed Glycosylation Enables the Stereocontrolled Synthesis of Selective SGLT2 Inhibitors for Type-2 Diabetes. <i>Chemistry - A European Journal</i> , 2018, 24, 2832-2836.	3.3	29
11	Development of Novel Melanocortin Receptor Agonists Based on the Cyclic Peptide Framework of Sunflower Trypsin Inhibitor-1. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 3674-3684.	6.4	29
12	Targeted delivery of antisense oligonucleotides to pancreatic β -cells. <i>Science Advances</i> , 2018, 4, eaat3386.	10.3	132
13	Exploring physicochemical space <i>via</i> a bioisostere of the trifluoromethyl and ethyl groups (BITE): attenuating lipophilicity in fluorinated analogues of Gilenya [®] for multiple sclerosis. <i>Chemical Communications</i> , 2018, 54, 12002-12005.	4.1	38
14	Discovery of a Series of Indole-2 Carboxamides as Selective Secreted Phospholipase A ₂ Type X (sPLA ₂ -X) Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 594-599.	2.8	9
15	Design of Selective sPLA ₂ -X Inhibitor (β)-2-[2-[Carbamoyl-6-(trifluoromethoxy)-1H-indol-1-yl]pyridine-2-yl]propanoic Acid. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 600-605.	2.8	5
16	Direct Synthesis of α -Alkyl Arylglycines by Organocatalytic Asymmetric Transfer Hydrogenation of α -Alkyl Aryl Imino Esters. <i>Organic Letters</i> , 2017, 19, 5541-5544.	4.6	14
17	Discovery of a novel pyrazole series of group X secreted phospholipase A2 inhibitor (sPLA2X) via fragment based virtual screening. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 5251-5255.	2.2	14