

Laurent Knerr

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4698876/publications.pdf>

Version: 2024-02-01

17
papers

470
citations

759233

12
h-index

839539

18
g-index

18
all docs

18
docs citations

18
times ranked

654
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|---|------|-----------|
| 1 | Targeted delivery of antisense oligonucleotides to pancreatic Î²-cells. <i>Science Advances</i> , 2018, 4, eaat3386. | 10.3 | 132 |
| 2 | Late-Stage Functionalization of Histidine in Unprotected Peptides. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 19096-19102. | 13.8 | 47 |
| 3 | 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 |
| 4 | 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 |
| 5 | Late-Stage Functionalization of Histidine in Unprotected Peptides. <i>Angewandte Chemie</i> , 2019, 131, 19272-19278. | 2.0 | 34 |
| 6 | Fluorine-Directed Glycosylation Enables the Stereocontrolled Synthesis of Selective SGLT2 Inhibitors for Type-II Diabetes. <i>Chemistry - A European Journal</i> , 2018, 24, 2832-2836. | 3.3 | 29 |
| 7 | 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 |
| 8 | Target-Directed Approaches for Screening Small Molecules against RNA Targets. <i>SLAS Discovery</i> , 2020, 25, 869-894. | 2.7 | 23 |
| 9 | Stereocontrolled Synthesis of Tetrafluoropentanol: Multivincinal Fluorinated Alkane Units for Drug Discovery. <i>Organic Letters</i> , 2019, 21, 7741-7745. | 4.6 | 20 |
| 10 | 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 |
| 11 | Direct Synthesis of <i>N</i> -Alkyl Arylglycines by Organocatalytic Asymmetric Transfer Hydrogenation of <i>N</i> -Alkyl Aryl Imino Esters. <i>Organic Letters</i> , 2017, 19, 5541-5544. | 4.6 | 14 |
| 12 | 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 |
| 13 | Iridium-Catalyzed Asymmetric Hydrogenation of <i>N</i> -Alkyl β -Aryl Furan-Containing Imines: an Efficient Route to Unnatural <i>N</i> -Alkyl Arylalanines and Related Derivatives.. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 578-584. | 4.3 | 12 |
| 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 | Melanocortin 1 Receptor Agonists Based on a Bivalent, Bicyclic Peptide Framework. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 9906-9915. | 6.4 | 6 |
| 16 | Design of Selective sPLA ₂ -X Inhibitor (β)-2-[2-[Carbamoyl-6-(trifluoromethoxy)-1 <i>H</i> -indol-1-yl]pyridine-2-yl]propanoic Acid. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 600-605. | 2.8 | 5 |
| 17 | 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 |