

Christoph Nitsche

List of Publications by Year in descending order

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Version: 2024-02-01

43
papers

2,783
citations

218677

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h-index

265206

42
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48
all docs

48
docs citations

48
times ranked

3574
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|------|-----------|
| 1 | Peptideâ€“Bismuth Bicycles: In Situ Access to Stable Constrained Peptides with Superior Bioactivity. <i>Angewandte Chemie - International Edition</i> , 2022, 61, . | 13.8 | 11 |
| 2 | Peptideâ€“Bismuth Bicycles: In Situ Access to Stable Constrained Peptides with Superior Bioactivity. <i>Angewandte Chemie</i> , 2022, 134, . | 2.0 | 3 |
| 3 | Genetic Encoding of Cyanopyridylalanine for Inâ€“Cell Protein Macrocyclization by the Nitrileâ€“Aminothioli Click Reaction. <i>Angewandte Chemie - International Edition</i> , 2022, 61, . | 13.8 | 15 |
| 4 | Organoarsenic probes to study proteins by NMR spectroscopy. <i>Chemical Communications</i> , 2022, 58, 701-704. | 4.1 | 1 |
| 5 | Paramagnetic Chemical Probes for Studying Biological Macromolecules. <i>Chemical Reviews</i> , 2022, 122, 9571-9642. | 47.7 | 36 |
| 6 | Antiviral cyclic peptides targeting the main protease of SARS-CoV-2. <i>Chemical Science</i> , 2022, 13, 3826-3836. | 7.4 | 29 |
| 7 | Main protease mutants of SARS-CoV-2 variants remain susceptible to nirmatrelvir. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2022, 62, 128629. | 2.2 | 131 |
| 8 | A biocompatible stapling reaction for <i>in situ</i> generation of constrained peptides. <i>Chemical Science</i> , 2021, 12, 669-674. | 7.4 | 25 |
| 9 | 2-Cyanoisonicotinamide Conjugation: A Facile Approach to Generate Potent Peptide Inhibitors of the Zika Virus Protease. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 732-737. | 2.8 | 21 |
| 10 | Challenges of short substrate analogues as SARS-CoV-2 main protease inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 50, 128333. | 2.2 | 26 |
| 11 | Targeting the protease of West Nile virus. <i>RSC Medicinal Chemistry</i> , 2021, 12, 1262-1272. | 3.9 | 6 |
| 12 | Nanoparticulate Inhibitors of Flavivirus Proteases from Zika, West Nile and Dengue Virus Are Cell-Permeable Antivirals. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 1955-1961. | 2.8 | 3 |
| 13 | Inhibitors of the Zika virus protease NS2B-NS3. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 126965. | 2.2 | 56 |
| 14 | Catching a Moving Target: Comparative Modeling of Flaviviral NS2B-NS3 Reveals Small Molecule Zika Protease Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 514-520. | 2.8 | 10 |
| 15 | The SARS-CoV-2 main protease as drug target. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127377. | 2.2 | 550 |
| 16 | Biocompatible Macrocyclization between Cysteine and 2-Cyanopyridine Generates Stable Peptide Inhibitors. <i>Organic Letters</i> , 2019, 21, 4709-4712. | 4.6 | 46 |
| 17 | Proteases from dengue, West Nile and Zika viruses as drug targets. <i>Biophysical Reviews</i> , 2019, 11, 157-165. | 3.2 | 51 |
| 18 | <i>De Novo</i> Discovery of Nonstandard Macrocyclic Peptides as Noncompetitive Inhibitors of the Zika Virus NS2B-NS3 Protease. <i>ACS Medicinal Chemistry Letters</i> , 2019, 10, 168-174. | 2.8 | 62 |

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|----|---|------|-----------|
| 19 | Trimethylsilyl tag for probing protein-ligand interactions by NMR. <i>Journal of Biomolecular NMR</i> , 2018, 70, 211-218. | 2.8 | 9 |
| 20 | NMR studies of ligand binding. <i>Current Opinion in Structural Biology</i> , 2018, 48, 16-22. | 5.7 | 48 |
| 21 | Strategies Towards Protease Inhibitors for Emerging Flaviviruses. <i>Advances in Experimental Medicine and Biology</i> , 2018, 1062, 175-186. | 1.6 | 33 |
| 22 | Small neutral Gd(III) tags for distance measurements in proteins by double electron-electron resonance experiments. <i>Physical Chemistry Chemical Physics</i> , 2018, 20, 23535-23545. | 2.8 | 22 |
| 23 | Intrinsic and Extrinsic Paramagnetic Probes. <i>New Developments in NMR</i> , 2018, , 42-84. | 0.1 | 9 |
| 24 | Peptide-Boronic Acid Inhibitors of Flaviviral Proteases: Medicinal Chemistry and Structural Biology. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 511-516. | 6.4 | 105 |
| 25 | Solution conformations of a linked construct of the Zika virus NS2B-NS3 protease. <i>Antiviral Research</i> , 2017, 142, 141-147. | 4.1 | 45 |
| 26 | Pseudocontact shifts in biomolecular NMR using paramagnetic metal tags. <i>Progress in Nuclear Magnetic Resonance Spectroscopy</i> , 2017, 98-99, 20-49. | 7.5 | 125 |
| 27 | Site-selective tagging of proteins by pnictogen-mediated self-assembly. <i>Chemical Communications</i> , 2017, 53, 10894-10897. | 4.1 | 15 |
| 28 | Crystal structure of Zika virus NS2B-NS3 protease in complex with a boronate inhibitor. <i>Science</i> , 2016, 353, 503-505. | 12.6 | 285 |
| 29 | The Medicinal Chemistry of Dengue Virus. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 5622-5649. | 6.4 | 114 |
| 30 | Sensitive NMR Approach for Determining the Binding Mode of Tightly Binding Ligand Molecules to Protein Targets. <i>Journal of the American Chemical Society</i> , 2016, 138, 4539-4546. | 13.7 | 53 |
| 31 | Dual inhibitors of the dengue and West Nile virus NS2B-NS3 proteases: Synthesis, biological evaluation and docking studies of novel peptide-hybrids. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 5748-5755. | 3.0 | 37 |
| 32 | Phenylalanine and Phenylglycine Analogues as Arginine Mimetics in Dengue Protease Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 7719-7733. | 6.4 | 69 |
| 33 | C-Terminal Residue Optimization and Fragment Merging: Discovery of a Potent Peptide-Hybrid Inhibitor of Dengue Protease. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 1037-1042. | 2.8 | 51 |
| 34 | Promiscuity and Selectivity in Covalent Enzyme Inhibition: A Systematic Study of Electrophilic Fragments. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 7590-7599. | 6.4 | 134 |
| 35 | Biochemistry and Medicinal Chemistry of the Dengue Virus Protease. <i>Chemical Reviews</i> , 2014, 114, 11348-11381. | 47.7 | 120 |
| 36 | The dengue virus NS2B-NS3 protease retains the closed conformation in the complex with BPTI. <i>FEBS Letters</i> , 2014, 588, 2206-2211. | 2.8 | 46 |

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|----|--|-----|-----------|
| 37 | Thiazolidinoneâ€“Peptide Hybrids as Dengue Virus Protease Inhibitors with Antiviral Activity in Cell Culture. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 8389-8403. | 6.4 | 110 |
| 38 | Cytotoxic betulin-derived hydroxypropargylamines trigger apoptosis. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 425-435. | 3.0 | 29 |
| 39 | Fluorimetric and HPLC-Based Dengue Virus Protease Assays Using a FRET Substrate. <i>Methods in Molecular Biology</i> , 2013, 1030, 221-236. | 0.9 | 19 |
| 40 | Aqueous microwave-assisted one-pot synthesis of N-substituted rhodanines. <i>Tetrahedron Letters</i> , 2012, 53, 5197-5201. | 1.4 | 42 |
| 41 | Retro peptide-hybrids as selective inhibitors of the Dengue virus NS2B-NS3 protease. <i>Antiviral Research</i> , 2012, 94, 72-79. | 4.1 | 78 |
| 42 | Arylcianoacrylamides as inhibitors of the Dengue and West Nile virus proteases. <i>Bioorganic and Medicinal Chemistry</i> , 2011, 19, 7318-7337. | 3.0 | 90 |
| 43 | Genetic Encoding of Cyanopyridylalanine for Inâ€“Cell Protein Macrocyclization by the Nitrileâ€“Amino-thiol Click Reaction. <i>Angewandte Chemie</i> , 0, , . | 2.0 | 0 |