Bruno Melillo

List of Publications by Year in descending order

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| # | Article | IF | CITATIONS |
|----|--|------|-----------|
| 1 | Crystallization-Based Synthetic Route to Antimalarial Agent BRD5018: Diazocene Ring Formation via a Staudinger-aza-Wittig Reaction on an Azetidine-Ribose Template. Organic Process Research and Development, 2022, 26, 817-831. | 1.3 | 5 |
| 2 | Bicyclic azetidines target acute and chronic stages of Toxoplasma gondii by inhibiting parasite phenylalanyl t-RNA synthetase. Nature Communications, 2022, 13, 459. | 5.8 | 7 |
| 3 | Inhibition of Plasmodium falciparum phenylalanine tRNA synthetase provides opportunity for antimalarial drug development. Structure, 2022, 30, 962-972.e3. | 1.6 | 4 |
| 4 | Stereochemical diversity as a source of discovery in chemical biology. Current Research in Chemical Biology, 2022, 2, 100028. | 1.4 | 21 |
| 5 | LPCAT3 Inhibitors Remodel the Polyunsaturated Phospholipid Content of Human Cells and Protect from Ferroptosis. ACS Chemical Biology, 2022, 17, 1607-1618. | 1.6 | 51 |
| 6 | Stabilizing the HIV-1 Envelope Glycoprotein State 2A Conformation. Journal of Virology, 2021, 95, . | 1.5 | 9 |
| 7 | Cryo-EM structures of HIV-1 trimer bound to CD4-mimetics BNM-III-170 and M48U1 adopt a CD4-bound open conformation. Nature Communications, 2021, 12, 1950. | 5.8 | 22 |
| 8 | Structural basis of malaria parasite phenylalanine tRNA-synthetase inhibition by bicyclic azetidines. Nature Communications, 2021, 12, 343. | 5.8 | 19 |
| 9 | Bicyclic azetidines kill the diarrheal pathogen <i>Cryptosporidium</i> in mice by inhibiting parasite phenylalanyl-tRNA synthetase. Science Translational Medicine, 2020, 12, . | 5.8 | 45 |
| 10 | An Activity-Guided Map of Electrophile-Cysteine Interactions in Primary Human T Cells. Cell, 2020, 182, 1009-1026.e29. | 13.5 | 194 |
| 11 | Elicitation of Cluster A and Co-Receptor Binding Site Antibodies Are Required to Eliminate HIV-1 Infected Cells. Microorganisms, 2020, 8, 710. | 1.6 | 7 |
| 12 | Ligandâ€Enabled βâ€Methylene C(sp 3)â^'H Arylation of Masked Aliphatic Alcohols. Angewandte Chemie, 2020, 132, 7857-7861. | 1.6 | 14 |
| 13 | Ligandâ€Enabled βâ€Methylene C(sp ³)â^'H Arylation of Masked Aliphatic Alcohols. Angewandte Chemie - International Edition, 2020, 59, 7783-7787. | 7.2 | 45 |
| 14 | Development of an Effective Scalable Enantioselective Synthesis of the HIV-1 Entry Inhibitor BNM-III-170 as the Bis-trifluoroacetate Salt. Organic Process Research and Development, 2019, 23, 2464-2469. | 1.3 | 21 |
| 15 | Strain-Dependent Activation and Inhibition of Human Immunodeficiency Virus Entry by a Specific PF-68742 Stereoisomer. Journal of Virology, 2019, 93, . | 1.5 | 1 |
| 16 | Modular, stereocontrolled C _β –H/C _α –C activation of alkyl carboxylic acids. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 8721-8727. | 3.3 | 39 |
| 17 | An Asymmetric Opening of HIV-1 Envelope Mediates Antibody-Dependent Cellular Cytotoxicity. Cell Host and Microbe, 2019, 25, 578-587.e5. | 5.1 | 93 |
| 18 | A Small-Molecule CD4-Mimetic Compound Protects Bone Marrow–Liver–Thymus Humanized Mice From HIV-1 Infection. Journal of Infectious Diseases, 2018, 218, 471-475. | 1.9 | 22 |

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|----|--|-----|-----------|
| 19 | SOSIP Changes Affect Human Immunodeficiency Virus Type 1 Envelope Glycoprotein Conformation and CD4 Engagement. Journal of Virology, 2018, 92, . | 1.5 | 24 |
| 20 | Synergistic Effects of Stereochemistry and Appendages on the Performance Diversity of a Collection of Synthetic Compounds. Journal of the American Chemical Society, 2018, 140, 11784-11790. | 6.6 | 47 |
| 21 | A CD4-mimetic compound enhances vaccine efficacy against stringent immunodeficiency virus challenge. Nature Communications, 2018, 9, 2363. | 5.8 | 46 |
| 22 | Activation and Inactivation of Primary Human Immunodeficiency Virus Envelope Glycoprotein Trimers by CD4-Mimetic Compounds. Journal of Virology, 2017, 91, . | 1.5 | 42 |
| 23 | BST-2 Expression Modulates Small CD4-Mimetic Sensitization of HIV-1-Infected Cells to Antibody-Dependent Cellular Cytotoxicity. Journal of Virology, 2017, 91, . | 1.5 | 40 |
| 24 | Residues in the gp41 Ectodomain Regulate HIV-1 Envelope Glycoprotein Conformational Transitions Induced by gp120-Directed Inhibitors. Journal of Virology, 2017, 91, . | 1.5 | 53 |
| 25 | Synthesis of a Bicyclic Azetidine with In Vivo Antimalarial Activity Enabled by Stereospecific, Directed C(sp ³)–H Arylation. Journal of the American Chemical Society, 2017, 139, 11300-11306. | 6.6 | 104 |
| 26 | Stereospecific Palladium-Catalyzed C–H Arylation of Pyroglutamic Acid Derivatives at the C3 Position Enabled by 8-Aminoquinoline as a Directing Group. Organic Letters, 2017, 19, 4424-4427. | 2.4 | 38 |
| 27 | Short Communication: Small-Molecule CD4 Mimetics Sensitize HIV-1-Infected Cells to Antibody-Dependent Cellular Cytotoxicity by Antibodies Elicited by Multiple Envelope Glycoprotein Immunogens in Nonhuman Primates. AIDS Research and Human Retroviruses, 2017, 33, 428-431. | 0.5 | 26 |
| 28 | Typeâ€II Anion Relay Chemistry: Exploiting Bifunctional Weinreb Amide Linchpins for the Oneâ€Pot Synthesis of Differentiated 1,3â€Diketones, Pyrans, and Spiroketals. Angewandte Chemie - International Edition, 2016, 55, 232-235. | 7.2 | 26 |
| 29 | Small CD4 Mimetics Prevent HIV-1 Uninfected Bystander CD4 + T Cell Killing Mediated by Antibody-dependent Cell-mediated Cytotoxicity. EBioMedicine, 2016, 3, 122-134. | 2.7 | 67 |
| 30 | Co-receptor Binding Site Antibodies Enable CD4-Mimetics to Expose Conserved Anti-cluster A ADCC Epitopes on HIV-1 Envelope Glycoproteins. EBioMedicine, 2016, 12, 208-218. | 2.7 | 65 |
| 31 | Computational Evaluation of HIV-1 gp120 Conformations of Soluble Trimeric gp140 Structures as Targets for de Novo Docking of First- and Second-Generation Small-Molecule CD4 Mimics. Journal of Chemical Information and Modeling, 2016, 56, 2069-2079. | 2.5 | 9 |
| 32 | Release of gp120 Restraints Leads to an Entry-Competent Intermediate State of the HIV-1 Envelope Glycoproteins. MBio, 2016, 7, . | 1.8 | 131 |
| 33 | Small-Molecule CD4-Mimics: Structure-Based Optimization of HIV-1 Entry Inhibition. ACS Medicinal Chemistry Letters, 2016, 7, 330-334. | 1.3 | 86 |
| 34 | Design, Synthesis, and Validation of an Effective, Reusable Silicon-Based Transfer Agent for Room-Temperature Pd-Catalyzed Cross-Coupling Reactions of Aryl and Heteroaryl Chlorides with Readily Available Aryl Lithium Reagents. Journal of the American Chemical Society, 2016, 138, 1836-1839. | 6.6 | 38 |
| 35 | Antibodies Elicited by Multiple Envelope Glycoprotein Immunogens in Primates Neutralize Primary Human Immunodeficiency Viruses (HIV-1) Sensitized by CD4-Mimetic Compounds. Journal of Virology, 2016, 90, 5031-5046. | 1.5 | 38 |
| 36 | An Effective Bifunctional Aldehyde Linchpin for Type II Anion Relay Chemistry: Development and Application to the Synthesis of a C16–C29 Fragment of Rhizopodin. Organic Letters, 2015, 17, 6242-6245. | 2.4 | 13 |

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| 37 | A Unified Synthetic Strategy to the Cryptocarya Family of Natural Products Exploiting Anion Relay Chemistry (ARC). Organic Letters, 2013, 15, 2282-2285. | 2.4 | 49 |