

Jenny Desantis

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3513326/publications.pdf>

Version: 2024-02-01

24
papers

763
citations

623188

14
h-index

610482

24
g-index

26
all docs

26
docs citations

26
times ranked

1047
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|-----------|
| 1 | Design and Synthesis of DiselenoBisBenzamides (DISEBAs) as Nucleocapsid Protein 7 (NCp7) Inhibitors with anti-HIV Activity. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 9601-9614. | 2.9 | 175 |
| 2 | A Broad Anti-influenza Hybrid Small Molecule That Potently Disrupts the Interaction of Polymerase Acidic Protein–Basic Protein 1 (PA-PB1) Subunits. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 3830-3842. | 2.9 | 81 |
| 3 | Understanding the Metabolism of Proteolysis Targeting Chimeras (PROTACs): The Next Step toward Pharmaceutical Applications. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 11615-11638. | 2.9 | 69 |
| 4 | Effects of MTX-23, a Novel PROTAC of Androgen Receptor Splice Variant-7 and Androgen Receptor, on CRPC Resistant to Second-Line Antiandrogen Therapy. <i>Molecular Cancer Therapeutics</i> , 2021, 20, 490-499. | 1.9 | 55 |
| 5 | Indomethacin-based PROTACs as pan-coronavirus antiviral agents. <i>European Journal of Medicinal Chemistry</i> , 2021, 226, 113814. | 2.6 | 46 |
| 6 | Polymerase Acidic Protein–Basic Protein 1 (PA–PB1) Protein–Protein Interaction as a Target for Next-Generation Anti-influenza Therapeutics. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 7699-7718. | 2.9 | 43 |
| 7 | Exploring the cycloheptathiophene-3-carboxamide scaffold to disrupt the interactions of the influenza polymerase subunits and obtain potent anti-influenza activity. <i>European Journal of Medicinal Chemistry</i> , 2017, 138, 128-139. | 2.6 | 38 |
| 8 | Efficient and regioselective one-step synthesis of 7-aryl-5-methyl- and 5-aryl-7-methyl-2-amino-[1,2,4]triazolo[1,5-a]pyrimidine derivatives. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 7944-7955. | 1.5 | 31 |
| 9 | Reducing Mutant Huntingtin Protein Expression in Living Cells by a Newly Identified RNA CAG Binder. <i>ACS Chemical Neuroscience</i> , 2018, 9, 1399-1408. | 1.7 | 29 |
| 10 | 4-(Phenoxy) and 4-(benzyloxy)benzamides as potent and selective inhibitors of mono-ADP-ribosyltransferase PARP10/ARTD10. <i>European Journal of Medicinal Chemistry</i> , 2018, 156, 93-102. | 2.6 | 23 |
| 11 | 1,2,4-Triazolo[1,5-a]pyrimidines as a Novel Class of Inhibitors of the HIV-1 Reverse Transcriptase-Associated Ribonuclease H Activity. <i>Molecules</i> , 2020, 25, 1183. | 1.7 | 23 |
| 12 | Potent and broad-spectrum cycloheptathiophene-3-carboxamide compounds that target the PA-PB1 interaction of influenza virus RNA polymerase and possess a high barrier to drug resistance. <i>Antiviral Research</i> , 2019, 165, 55-64. | 1.9 | 20 |
| 13 | Inhibition of Influenza Virus Polymerase by Interfering with Its Protein–Protein Interactions. <i>ACS Infectious Diseases</i> , 2021, 7, 1332-1350. | 1.8 | 18 |
| 14 | 2-Phenylquinazolinones as dual-activity tankyrase-kinase inhibitors. <i>Scientific Reports</i> , 2018, 8, 1680. | 1.6 | 16 |
| 15 | From cycloheptathiophene-3-carboxamide to oxazinone-based derivatives as allosteric HIV-1 ribonuclease H inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019, 34, 55-74. | 2.5 | 16 |
| 16 | Studies on Cycloheptathiophene-3-carboxamide Derivatives as Allosteric HIV-1 Ribonuclease...H Inhibitors. <i>ChemMedChem</i> , 2016, 11, 1709-1720. | 1.6 | 15 |
| 17 | Proteolysis targeting chimeras in antiviral research. <i>Future Medicinal Chemistry</i> , 2022, 14, 459-462. | 1.1 | 14 |
| 18 | Recent advances in the identification of Tat-mediated transactivation inhibitors: progressing toward a functional cure of HIV. <i>Future Medicinal Chemistry</i> , 2016, 8, 421-442. | 1.1 | 12 |

| # | ARTICLE | IF | CITATIONS |
|----|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|-----------|
| 19 | Antitubercular polyhalogenated phenothiazines and phenoselenazine with reduced binding to CNS receptors. <i>European Journal of Medicinal Chemistry</i> , 2020, 201, 112420. | 2.6 | 12 |
| 20 | Discovery of novel SARS-CoV-2 inhibitors targeting the main protease Mpro by virtual screenings and hit optimization. <i>Antiviral Research</i> , 2022, 204, 105350. | 1.9 | 11 |
| 21 | Automatic Identification of Lansoprazole Degradants under Stress Conditions by LC-HRMS with MassChemSite and WebChembase. <i>Journal of Chemical Information and Modeling</i> , 2021, 61, 2706-2719. | 2.5 | 5 |
| 22 | CROMATIC: <i>Cross-Relationship Mapping of Cavitities from Coronavirus</i> . <i>Journal of Chemical Information and Modeling</i> , 0, , . | 2.5 | 4 |
| 23 | An overview on small molecules acting as broad-spectrum agents for yellow fever infection. <i>Expert Opinion on Drug Discovery</i> , 2022, 17, 755-773. | 2.5 | 3 |
| 24 | Design and Synthesis of WM5 Analogues as HIV-1 TAR RNA Binders. <i>Open Medicinal Chemistry Journal</i> , 2019, 13, 16-28. | 0.9 | 2 |