Giovanna Zinzalla

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2936320/publications.pdf

Version: 2024-02-01

34 1,061 17
papers citations h-index

17 30
h-index g-index

48 48 all docs docs citations

48 times ranked 1870 citing authors

| # | Article | IF | CITATIONS |
|----|--|-------------|-----------|
| 1 | Targeting protein–protein interactions for therapeutic intervention: a challenge for the future. Future Medicinal Chemistry, 2009, 1, 65-93. | 2.3 | 221 |
| 2 | Chemical variation of natural product-like scaffolds: design and synthesis of spiroketal derivatives. Organic and Biomolecular Chemistry, 2006, 4, 1977. | 2.8 | 85 |
| 3 | A selective high affinity MYC-binding compound inhibits MYC:MAX interaction and MYC-dependent tumor cell proliferation. Scientific Reports, 2018, 8, 10064. | 3.3 | 85 |
| 4 | Synthesis of Chiral Chromium Tricarbonyl Labeled Thymine PNA Monomers via the Ugi Reaction. Organic Letters, 2002, 4, 4341-4344. | 4.6 | 61 |
| 5 | Observation of unphosphorylated STAT3 core protein binding to target ⟨i>ds⟨/i>DNA by PEMSA and Xâ€ray crystallography. FEBS Letters, 2013, 587, 833-839. | 2.8 | 60 |
| 6 | Small-Molecule Inhibition of c-MYC:MAX Leucine Zipper Formation Is Revealed by Ion Mobility Mass Spectrometry. Journal of the American Chemical Society, 2012, 134, 19384-19392. | 13.7 | 53 |
| 7 | Chemical Variation of Natural-Product-Like Scaffolds: Design, Synthesis, and Biological Activity of Fused Bicyclic Acetal Derivatives. Angewandte Chemie - International Edition, 2007, 46, 2493-2496. | 13.8 | 51 |
| 8 | Investigation of the protein alkylation sites of the STAT3:STAT3 inhibitor Stattic by mass spectrometry. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 4719-4722. | 2.2 | 45 |
| 9 | Crystal Structures and Nuclear Magnetic Resonance Studies of the Apo Form of the c-MYC:MAX bHLHZip Complex Reveal a Helical Basic Region in the Absence of DNA. Biochemistry, 2019, 58, 3144-3154. | 2.5 | 38 |
| 10 | A new ferrocene conjugate of a tyrosine PNA monomer: synthesis and electrochemical properties. Journal of Organometallic Chemistry, 2004, 689, 4791-4802. | 1.8 | 35 |
| 11 | Naturalâ€Productâ€Like Spiroketals and Fused Bicyclic Acetals as Potential Therapeutic Agents for B ell Chronic Lymphocytic Leukaemia. ChemMedChem, 2008, 3, 1922-1935. | 3. 2 | 34 |
| 12 | The SWI/SNF Subunit INI1 Contains an N-Terminal Winged Helix DNA Binding Domain that Is a Target for Mutations in Schwannomatosis. Structure, 2015, 23, 1344-1349. | 3.3 | 33 |
| 13 | Facile nucleophilic substitution at the C3a tertiary carbon of the 3a-bromohexahydropyrrolo[2,3-b]indole scaffold. Organic and Biomolecular Chemistry, 2010, 8, 5294. | 2.8 | 28 |
| 14 | Targeting protein–protein interactions (PPIs) of transcription factors: Challenges of intrinsically disordered proteins (IDPs) and regions (IDRs). Progress in Biophysics and Molecular Biology, 2015, 119, 41-46. | 2.9 | 27 |
| 15 | The structure of <scp>INI</scp> 1/ <scp>hSNF</scp> 5 <scp>RPT</scp> 1 and its interactions with the câ€ <scp>MYC</scp> : <scp>MAX</scp> heterodimer provide insights into the interplay between <scp>MYC</scp> and the <scp>SWI</scp> / <scp>SNF</scp> chromatin remodeling complex. FEBS lournal, 2018, 285, 4165-4180. | 4.7 | 22 |
| 16 | Molecular Dynamics Studies of the STAT3 Homodimer:DNA Complex: Relationships between STAT3 Mutations and Protein–DNA Recognition. Journal of Chemical Information and Modeling, 2012, 52, 1179-1192. | 5.4 | 21 |
| 17 | A New Way Forward in Cancer Drug Discovery: Inhibiting the SWI/SNF Chromatin Remodelling Complex. ChemBioChem, 2016, 17, 677-682. | 2.6 | 21 |
| 18 | Facile oxidation of electron-poor benzo[b]thiophenes to the corresponding sulfones with an aqueous solution of H2O2 and P2O5. Chemical Communications, 2010, 46, 2289. | 4.1 | 20 |

| # | Article | IF | CITATIONS |
|----|--|-----|-----------|
| 19 | Stereoselective hetero Diels–Alder reactions of chiral tricarbonyl (η6-benzaldehyde)chromium complexes. Tetrahedron: Asymmetry, 2001, 12, 2159-2167. | 1.8 | 18 |
| 20 | A Thymine-PNA Monomer as New Isocyanide Component in the Ugi Reaction: A Direct Entry to PNA Dimers. Synlett, 2004, 2004, 1044-1048. | 1.8 | 18 |
| 21 | Structure of the BRK domain of the SWI/SNF chromatin remodeling complex subunit BRG1 reveals a potential role in protein–protein interactions. Protein Science, 2020, 29, 1033-1039. | 7.6 | 17 |
| 22 | A novel small-molecule inhibitor of IL-6 signalling. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 7029-7032. | 2.2 | 16 |
| 23 | Polymer-Supported Haloarene Chromium Dicarbonyl Isonitrile Complexes:  A Study of Their Synthesis and Reactivity. ACS Combinatorial Science, 2003, 5, 809-813. | 3.3 | 12 |
| 24 | Paving the way to targeting HECT ubiquitin ligases. Future Medicinal Chemistry, 2015, 7, 2107-2111. | 2.3 | 6 |
| 25 | Targeting MYC: is it getting any easier?. Future Medicinal Chemistry, 2016, 8, 1899-1902. | 2.3 | 6 |
| 26 | SWI/SNF subunit BAF155 N-terminus structure informs the impact of cancer-associated mutations and reveals a potential drug binding site. Communications Biology, 2021, 4, 528. | 4.4 | 5 |
| 27 | Tetracycline analogues with a selective inhibitory effect on HIF-1α. MedChemComm, 2014, 5, 923. | 3.4 | 3 |
| 28 | One-Pot Synthesis of Fused-Tetracyclic Scaffolds Employing a Lewis Acid Promoted Domino Reaction of Naphthoquinones. Synthesis, 2011, 2011, 2321-2333. | 2.3 | 2 |
| 29 | Biophysical and Structural Methods to Study the bHLHZip Region of Human c-MYC. Methods in Molecular Biology, 2021, 2318, 21-43. | 0.9 | 1 |
| 30 | 113 Novel small-molecule inhibitors of Interleukin-6 (IL-6) signalling. European Journal of Cancer, Supplement, 2010, 8, 43. | 2.2 | 0 |
| 31 | Abstract 5454: Novel STAT3:STAT3 small-molecule inhibitors as potential anticancer agents. , 2010, , . | | 0 |
| 32 | Abstract 1382: Use of polarized light spectroscopy (CD) to study STAT3 folding and STAT3:ligand interactions., 2011,,. | | 0 |
| 33 | Abstract 279: Use of GFP-STAT3 \hat{l}^2 tc for EMSA analysis of protein-protein and protein-DNA interactions in tumorigenic signalling pathways. , 2011, , . | | 0 |
| 34 | Abstract 3952: Selective high affinity MYC-binding compound inhibits MYC-MAX interaction and MYC-dependent tumor cell growth. , 2018, , . | | 0 |