Sandeep Rana

List of Publications by Year in descending order

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| # | Article | IF | CITATIONS |
|----|---|------|-----------|
| 1 | Stapling proteins in the RELA complex inhibits TNFα-induced nuclear translocation of RELA. RSC Chemical Biology, 2022, 3, 32-36. | 4.1 | 6 |
| 2 | Spirocyclic dimer SpiD7 activates the unfolded protein response to selectively inhibit growth and induce apoptosis of cancer cells. Journal of Biological Chemistry, 2022, 298, 101890. | 3.4 | 5 |
| 3 | Dimers of isatin derived α-methylene-γ-butyrolactone as potent anti-cancer agents. Bioorganic and Medicinal Chemistry Letters, 2022, 65, 128713. | 2.2 | 5 |
| 4 | Small-molecule IKKβ activation modulator (IKAM) targets MAP3K1 and inhibits pancreatic tumor growth. Proceedings of the National Academy of Sciences of the United States of America, 2022, 119, e2115071119. | 7.1 | 3 |
| 5 | Selective killing of homologous recombination-deficient cancer cell lines by inhibitors of the RPA:RAD52 protein-protein interaction. PLoS ONE, 2021, 16, e0248941. | 2.5 | 13 |
| 6 | Small molecule induced polymerization of BCL6 facilitates SIAH1 mediated degradation. Signal Transduction and Targeted Therapy, 2021, 6, 142. | 17.1 | 3 |
| 7 | PIK3C3 Inhibition Promotes Sensitivity to Colon Cancer Therapy by Inhibiting Cancer Stem Cells. Cancers, 2021, 13, 2168. | 3.7 | 28 |
| 8 | Aminopyrazole based CDK9 PROTAC sensitizes pancreatic cancer cells to venetoclax. Bioorganic and Medicinal Chemistry Letters, 2021, 43, 128061. | 2.2 | 30 |
| 9 | Inhibitors, PROTACs and Molecular Glues as Diverse Therapeutic Modalities to Target Cyclin-Dependent Kinase. Cancers, 2021, 13, 5506. | 3.7 | 17 |
| 10 | A Novel Spirocyclic Dimer (36-286) Targeting the NF-Kappa B Pathway Displays Potent Anti-Tumor Properties in Chronic Lymphocytic Leukemia. Blood, 2021, 138, 1186-1186. | 1.4 | 0 |
| 11 | EHD1 and RUSC2 Control Basal Epidermal Growth Factor Receptor Cell Surface Expression and Recycling. Molecular and Cellular Biology, 2020, 40, . | 2.3 | 8 |
| 12 | Synthesis and SAR studies of novel 1,2,4-oxadiazole-sulfonamide based compounds as potential anticancer agents for colorectal cancer therapy. Bioorganic Chemistry, 2020, 98, 103754. | 4.1 | 29 |
| 13 | Symbiotic prodrugs (SymProDs) dual targeting of NFkappaB and CDK. Chemical Biology and Drug Design, 2020, 96, 773-784. | 3.2 | 10 |
| 14 | Synthesis, Anticancer Evaluation and DNAâ€Binding Spectroscopic Insights of Quinolineâ€Based 1,3,4â€Oxadiazoleâ€1,2,3â€triazole Conjugates. ChemistrySelect, 2019, 4, 12176-12182. | 1.5 | 19 |
| 15 | A mitotic CDK5-PP4 phospho-signaling cascade primes 53BP1 for DNA repair in G1. Nature Communications, 2019, 10, 4252. | 12.8 | 17 |
| 16 | Selective degradation of CDK6 by a palbociclib based PROTAC. Bioorganic and Medicinal Chemistry Letters, 2019, 29, 1375-1379. | 2.2 | 95 |
| 17 | CDK5 Inhibitor Downregulates Mcl-1 and Sensitizes Pancreatic Cancer Cell Lines to Navitoclax. Molecular Pharmacology, 2019, 96, 419-429. | 2.3 | 21 |
| 18 | Chemical Genetic Screens Identify Kinase Inhibitor Combinations that Target Anti-Apoptotic Proteins for Cancer Therapy. ACS Chemical Biology, 2018, 13, 1148-1152. | 3.4 | 10 |

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|----|--|-----|-----------|
| 19 | Synthesis of aminopyrazole analogs and their evaluation as CDK inhibitors for cancer therapy. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 3736-3740. | 2.2 | 16 |
| 20 | Recent Advances in Cancer Drug Development: Targeting Induced Myeloid Cell Leukemia-1 (Mcl-1) Differentiation Protein. Current Medicinal Chemistry, 2018, 24, 4488-4514. | 2.4 | 13 |
| 21 | Characterization of CDK(5) inhibitor, 20-223 (aka CP668863) for colorectal cancer therapy. Oncotarget, 2018, 9, 5216-5232. | 1.8 | 22 |
| 22 | Development of 1-((1,4- <i>trans</i>)-4-Aryloxycyclohexyl)-3-arylurea Activators of Heme-Regulated Inhibitor as Selective Activators of the Eukaryotic Initiation Factor 2 Alpha (eIF2α) Phosphorylation Arm of the Integrated Endoplasmic Reticulum Stress Response. Journal of Medicinal Chemistry, 2017, 60, 5392-5406. | 6.4 | 17 |
| 23 | Chemically induced degradation of CDK9 by a proteolysis targeting chimera (PROTAC). Chemical Communications, 2017, 53, 7577-7580. | 4.1 | 167 |
| 24 | Cyclin Dependent Kinase 9 Inhibitors for Cancer Therapy. Journal of Medicinal Chemistry, 2016, 59, 8667-8684. | 6.4 | 121 |
| 25 | A quinoxaline urea analog uncouples inflammatory and pro-survival functions of IKKβ. Immunology Letters, 2015, 168, 319-324. | 2.5 | 5 |
| 26 | Micellar formulation of indocyanine green for phototherapy of melanoma. Journal of Controlled Release, 2015, 220, 130-140. | 9.9 | 49 |
| 27 | Small Molecule Adenosine 5′-Monophosphate Activated Protein Kinase (AMPK) Modulators and Human Diseases. Journal of Medicinal Chemistry, 2015, 58, 2-29. | 6.4 | 51 |
| 28 | Face selective reduction of the exocyclic double bond in isatin derived spirocyclic lactones. Organic and Biomolecular Chemistry, 2013, 11, 244-247. | 2.8 | 32 |