Wu Zhong

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2651477/publications.pdf Version: 2024-02-01

| | | 393982 | 253896 |
|----------|----------------|--------------|----------------|
| 42 | 11,087 | 19 | 43 |
| papers | citations | h-index | g-index |
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| 43 | 43 | 43 | 19475 |
| all docs | docs citations | times ranked | citing authors |
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| # | Article | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | Remdesivir and chloroquine effectively inhibit the recently emerged novel coronavirus (2019-nCoV) in vitro. Cell Research, 2020, 30, 269-271. | 5.7 | 5,527 |
| 2 | Analysis of therapeutic targets for SARS-CoV-2 and discovery of potential drugs by computational methods. Acta Pharmaceutica Sinica B, 2020, 10, 766-788. | 5.7 | 1,704 |
| 3 | Hydroxychloroquine, a less toxic derivative of chloroquine, is effective in inhibiting SARS-CoV-2 infection in vitro. Cell Discovery, 2020, 6, 16. | 3.1 | 1,643 |
| 4 | Experimental Treatment with Favipiravir for COVID-19: An Open-Label Control Study. Engineering, 2020, 6, 1192-1198. | 3.2 | 989 |
| 5 | Mortality outcomes with hydroxychloroquine and chloroquine in COVID-19 from an international collaborative meta-analysis of randomized trials. Nature Communications, 2021, 12, 2349. | 5.8 | 194 |
| 6 | Pathological features of COVID-19-associated lung injury: a preliminary proteomics report based on clinical samples. Signal Transduction and Targeted Therapy, 2020, 5, 240. | 7.1 | 148 |
| 7 | Anti-SARS-CoV-2 Potential of Artemisinins In Vitro. ACS Infectious Diseases, 2020, 6, 2524-2531. | 1.8 | 117 |
| 8 | Antibody–drug conjugates: Recent advances in linker chemistry. Acta Pharmaceutica Sinica B, 2021, 11, 3889-3907. | 5.7 | 114 |
| 9 | Host Calcium Channels and Pumps in Viral Infections. Cells, 2020, 9, 94. | 1.8 | 104 |
| 10 | Activation of the MAPK11/12/13/14 (p38 MAPK) pathway regulates the transcription of autophagy genes in response to oxidative stress induced by a novel copper complex in HeLa cells. Autophagy, 2014, 10, 1285-1300. | 4.3 | 82 |
| 11 | De Novo Design of α-Helical Lipopeptides Targeting Viral Fusion Proteins: A Promising Strategy for Relatively Broad-Spectrum Antiviral Drug Discovery. Journal of Medicinal Chemistry, 2018, 61, 8734-8745. | 2.9 | 41 |
| 12 | Design and synthesis of piperidine derivatives as novel human heat shock protein 70 inhibitors for the treatment of drug-resistant tumors. European Journal of Medicinal Chemistry, 2015, 97, 19-31. | 2.6 | 30 |
| 13 | Development and Properties of Valine-Alanine based Antibody-Drug Conjugates with Monomethyl Auristatin E as the Potent Payload. International Journal of Molecular Sciences, 2017, 18, 1860. | 1.8 | 30 |
| 14 | Small Molecule Inhibitor of ATPase Activity of HSP70 as a Broad-Spectrum Inhibitor against Flavivirus Infections. ACS Infectious Diseases, 2020, 6, 832-843. | 1.8 | 28 |
| 15 | Improvement of the <i>C</i> -glycosylation Step for the Synthesis of Remdesivir. Organic Process Research and Development, 2020, 24, 1772-1777. | 1.3 | 26 |
| 16 | Novel Silyl Ether-Based Acid-Cleavable Antibody-MMAE Conjugates with Appropriate Stability and Efficacy. Cancers, 2019, 11, 957. | 1.7 | 25 |
| 17 | Comparative Antiviral Efficacy of Viral Protease Inhibitors against the Novel SARS-CoV-2 In Vitro. Virologica Sinica, 2020, 35, 776-784. | 1.2 | 24 |
| 18 | Research progress on repositioning drugs and specific therapeutic drugs for SARS-CoV-2. Future Medicinal Chemistry, 2020, 12, 1565-1578. | 1,1 | 22 |

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|----|--|-----|-----------|
| 19 | Antibody-Drug Conjugate Using Ionized Cys-Linker-MMAE as the Potent Payload Shows Optimal Therapeutic Safety. Cancers, 2020, 12, 744. | 1.7 | 22 |
| 20 | Sera proteomic features of active and recovered COVID-19 patients: potential diagnostic and prognostic biomarkers. Signal Transduction and Targeted Therapy, 2021, 6, 216. | 7.1 | 22 |
| 21 | Pathological features of COVID-19-associated liver injury—a preliminary proteomics report based on clinical samples. Signal Transduction and Targeted Therapy, 2021, 6, 9. | 7.1 | 17 |
| 22 | Development of Novel Anti-influenza Thiazolides with Relatively Broad-Spectrum Antiviral Potentials. Antimicrobial Agents and Chemotherapy, 2020, 64, . | 1.4 | 16 |
| 23 | A bifunctional molecule-based strategy for the development of theranostic antibody-drug conjugate. Theranostics, 2021, 11, 2550-2563. | 4.6 | 15 |
| 24 | Application of omics technology to combat the COVIDâ€19 pandemic. MedComm, 2021, 2, 381-401. | 3.1 | 11 |
| 25 | Development of bifunctional anti-PD-L1 antibody MMAE conjugate with cytotoxicity and immunostimulation. Bioorganic Chemistry, 2021, 116, 105366. | 2.0 | 11 |
| 26 | Ebola virus VP35 hijacks the PKA-CREB1 pathway for replication and pathogenesis by AKIP1 association. Nature Communications, 2022, 13, 2256. | 5.8 | 11 |
| 27 | Molnupiravir and Its Active Form, EIDD-1931, Show Potent Antiviral Activity against Enterovirus Infections In Vitro and In Vivo. Viruses, 2022, 14, 1142. | 1.5 | 10 |
| 28 | Design, synthesis and pharmacological evaluation of a novel mTOR-targeted anti-EV71 agent. European Journal of Medicinal Chemistry, 2019, 175, 172-186. | 2.6 | 9 |
| 29 | Nafamostat mesylate as a broad-spectrum candidate for the treatment of flavivirus infections by targeting envelope proteins. Antiviral Research, 2022, 202, 105325. | 1.9 | 9 |
| 30 | Rapid Neutralization Testing System for Zika Virus Based on an Enzyme-Linked Immunospot Assay. ACS Infectious Diseases, 2020, 6, 811-819. | 1.8 | 8 |
| 31 | Novel antibody-drug conjugate with UV-controlled cleavage mechanism for cytotoxin release. Bioorganic Chemistry, 2021, 111, 104475. | 2.0 | 8 |
| 32 | LDL receptor-related protein 1 (LRP1), a novel target for opening the blood-labyrinth barrier (BLB). Signal Transduction and Targeted Therapy, 2022, 7, . | 7.1 | 8 |
| 33 | Development of a Novel Dual-Order Protein-Based Nanodelivery Carrier That Rapidly Targets Low-Grade Gliomas with Microscopic Metastasis <i>in Vivo</i> . ACS Omega, 2020, 5, 20653-20663. | 1.6 | 7 |
| 34 | Synthesis and evaluation of highly releasable and structurally stable antibody-SN-38-conjugates. Drug Delivery, 2021, 28, 2603-2617. | 2.5 | 7 |
| 35 | From prodrug to pro-prodrug: hypoxia-sensitive antibody–drug conjugates. Signal Transduction and Targeted Therapy, 2022, 7, 20. | 7.1 | 7 |
| 36 | In vitro and in vivo antiviral activity of Maqian (Zanthoxylum myriacanthum var. pubescens) essential oil and its major constituents against strains of influenza virus. Industrial Crops and Products, 2022, 177, 114524. | 2.5 | 6 |

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|----|--|-----|-----------|
| 37 | Design, synthesis and biological evaluation of 2-hydrazinyladenosine derivatives as A2A adenosine receptor ligands. European Journal of Medicinal Chemistry, 2019, 179, 310-324. | 2.6 | 4 |
| 38 | Tilorone confers robust in vitro and in vivo antiviral effects against severe fever with thrombocytopenia syndrome virus. Virologica Sinica, 2022, 37, 145-148. | 1.2 | 4 |
| 39 | The CDK1 inhibitor, Ro-3306, is a potential antiviral candidate against influenza virus infection. Antiviral Research, 2022, 201, 105296. | 1.9 | 4 |
| 40 | Design, synthesis and biological activity evaluation of a series of bardoxolone methyl prodrugs. Bioorganic Chemistry, 2022, 124, 105831. | 2.0 | 3 |
| 41 | Azelnidipine Exhibits In Vitro and In Vivo Antiviral Effects against Flavivirus Infections by Targeting the Viral RdRp. Viruses, 2022, 14, 1228. | 1.5 | 3 |
| 42 | Development of applicable thiol-linked antibody–drug conjugates with improved stability and therapeutic index. Drug Delivery, 2022, 29, 754-766. | 2.5 | 2 |