

Ashley N Brown

List of Publications by Year in descending order

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Version: 2024-02-01

24
papers

668
citations

687363

13
h-index

642732

23
g-index

24
all docs

24
docs citations

24
times ranked

1145
citing authors

#	ARTICLE	IF	CITATIONS
1	A sensitive electrochemical immunosensor for label-free detection of Zika-virus protein. Scientific Reports, 2018, 8, 9700.	3.3	148
2	Modeling the viral dynamics of SARS-CoV-2 infection. Mathematical Biosciences, 2020, 328, 108438.	1.9	120
3	Preclinical Evaluations To Identify Optimal Linezolid Regimens for Tuberculosis Therapy. MBio, 2015, 6, e01741-15.	4.1	60
4	Chikungunya Virus: In Vitro Response to Combination Therapy With Ribavirin and Interferon Alfa 2a. Journal of Infectious Diseases, 2016, 214, 1192-1197.	4.0	45
5	The effectiveness of antiviral agents with broad-spectrum activity against chikungunya virus varies between host cell lines. Antiviral Chemistry and Chemotherapy, 2018, 26, 204020661880758.	0.6	37
6	Oseltamivir-zanamivir combination therapy suppresses drug-resistant H1N1 influenza A viruses in the hollow fiber infection model (HFIM) system. European Journal of Pharmaceutical Sciences, 2018, 111, 443-449.	4.0	34
7	Zika Virus Replication Is Substantially Inhibited by Novel Favipiravir and Interferon Alpha Combination Regimens. Antimicrobial Agents and Chemotherapy, 2018, 62, .	3.2	29
8	Effect of Half-Life on the Pharmacodynamic Index of Zanamivir against Influenza Virus Delineated by a Mathematical Model. Antimicrobial Agents and Chemotherapy, 2011, 55, 1747-1753.	3.2	27
9	<i>In Vitro</i> System for Modeling Influenza A Virus Resistance under Drug Pressure. Antimicrobial Agents and Chemotherapy, 2010, 54, 3442-3450.	3.2	25
10	Zanamivir, at 600 Milligrams Twice Daily, Inhibits Oseltamivir-Resistant 2009 Pandemic H1N1 Influenza Virus in an <i>In Vitro</i> Hollow-Fiber Infection Model System. Antimicrobial Agents and Chemotherapy, 2011, 55, 1740-1746.	3.2	24
11	Clinical Regimens of Favipiravir Inhibit Zika Virus Replication in the Hollow-Fiber Infection Model. Antimicrobial Agents and Chemotherapy, 2018, 62, .	3.2	19
12	Antiviral Effects of Clinically-Relevant Interferon- α and Ribavirin Regimens against Dengue Virus in the Hollow Fiber Infection Model (HFIM). Viruses, 2018, 10, 317.	3.3	18
13	Pharmacokinetic Determinants of Virological Response to Raltegravir in the <i>In Vitro</i> Pharmacodynamic Hollow-Fiber Infection Model System. Antimicrobial Agents and Chemotherapy, 2015, 59, 3771-3777.	3.2	13
14	Application of pharmacometrics and quantitative systems pharmacology to cancer therapy: The example of luminal a breast cancer. Pharmacological Research, 2017, 124, 20-33.	7.1	13
15	Pharmacodynamic Analysis of a Serine Protease Inhibitor, MK-4519, against Hepatitis C Virus Using a Novel <i>In Vitro</i> Pharmacodynamic System. Antimicrobial Agents and Chemotherapy, 2012, 56, 1170-1181.	3.2	12
16	Utility of a Novel Three-Dimensional and Dynamic (3DD) Cell Culture System for PK/PD Studies: Evaluation of a Triple Combination Therapy at Overcoming Anti-HER2 Treatment Resistance in Breast Cancer. Frontiers in Pharmacology, 2018, 9, 403.	3.5	8
17	Combination Regimens of Favipiravir Plus Interferon Alpha Inhibit Chikungunya Virus Replication in Clinically Relevant Human Cell Lines. Microorganisms, 2021, 9, 307.	3.6	7
18	Antiviral Evaluation of UV-4B and Interferon-Alpha Combination Regimens against Dengue Virus. Viruses, 2021, 13, 771.	3.3	7

#	ARTICLE	IF	CITATIONS
19	Sofosbuvir (SOF) Suppresses Ledipasvir (LDV)-resistant Mutants during SOF/LDV Combination Therapy against Genotype 1b Hepatitis C Virus (HCV). <i>Scientific Reports</i> , 2017, 7, 14421.	3.3	6
20	Searching for synergy: Identifying optimal antiviral combination therapy using Hepatitis C virus (HCV) agents in a replicon system. <i>Antiviral Research</i> , 2017, 146, 149-152.	4.1	5
21	Antiviral Activity of the Propylamylatin TM Formula against the Novel Coronavirus SARS-CoV-2 In Vitro Using Direct Injection and Gas Assays in Virus Suspensions. <i>Viruses</i> , 2021, 13, 415.	3.3	5
22	UV-4B potently inhibits replication of multiple SARS-CoV-2 strains in clinically relevant human cell lines. <i>Frontiers in Bioscience</i> , 2022, 27, 1.	2.1	4
23	1088. A Whole-Body Quantitative System Pharmacology Physiologically-Based Pharmacokinetic (QSP/PBPK) Model to Support Dose Selection of ADG20: an Extended Half-Life Monoclonal Antibody Being Developed for the Treatment of Coronavirus Disease (COVID-19). <i>Open Forum Infectious Diseases</i> , 2021, 8, S635-S635.	0.9	2
24	Reply to Scagnolari et al.. <i>Journal of Infectious Diseases</i> , 2016, 215, jiw580.	4.0	0