

Yingshan Han

List of Publications by Year in descending order

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Version: 2024-02-01

13
papers

391
citations

840585

11
h-index

1125617

13
g-index

14
all docs

14
docs citations

14
times ranked

798
citing authors

#	ARTICLE	IF	CITATIONS
1	Antimalarial drugs and their metabolites are potent Zika virus inhibitors. <i>Journal of Medical Virology</i> , 2019, 91, 1182-1190.	2.5	36
2	The S230R Integrase Substitution Associated With Virus Load Rebound During Dolutegravir Monotherapy Confers Low-Level Resistance to Integrase Strand-Transfer Inhibitors. <i>Journal of Infectious Diseases</i> , 2018, 218, 698-706.	1.9	40
3	The antimalarial drug amodiaquine possesses anti-ZIKA virus activities. <i>Journal of Medical Virology</i> , 2018, 90, 796-802.	2.5	43
4	Investigational drugs for the treatment of Zika virus infection: a preclinical and clinical update. <i>Expert Opinion on Investigational Drugs</i> , 2018, 27, 951-962.	1.9	20
5	Purification of Zika virus RNA-dependent RNA polymerase and its use to identify small-molecule Zika inhibitors. <i>Journal of Antimicrobial Chemotherapy</i> , 2017, 72, dkw514.	1.3	55
6	JAK-STAT Signaling Pathways and Inhibitors Affect Reversion of Envelope-Mutated HIV-1. <i>Journal of Virology</i> , 2017, 91, .	1.5	11
7	The R263K Dolutegravir Resistance-Associated Substitution Progressively Decreases HIV-1 Integration. <i>MBio</i> , 2017, 8, .	1.8	14
8	Investigational HIV integrase inhibitors in phase I and phase II clinical trials. <i>Expert Opinion on Investigational Drugs</i> , 2017, 26, 1207-1213.	1.9	16
9	Evaluation of Sofosbuvir (2'-D-2-deoxy-2-fluoro-2'-C-methyluridine) as an inhibitor of Dengue virus replication #. <i>Scientific Reports</i> , 2017, 7, 6345.	1.6	58
10	Identification of a Pyridoxine-Derived Small-Molecule Inhibitor Targeting Dengue Virus RNA-Dependent RNA Polymerase. <i>Antimicrobial Agents and Chemotherapy</i> , 2016, 60, 600-608.	1.4	33
11	The R263K mutation in HIV integrase that is selected by dolutegravir may actually prevent clinically relevant resistance to this compound. <i>Journal of the International AIDS Society</i> , 2014, 17, 19518.	1.2	10
12	HIV-1 Group O Integrase Displays Lower Enzymatic Efficiency and Higher Susceptibility to Raltegravir than HIV-1 Group M Subtype B Integrase. <i>Antimicrobial Agents and Chemotherapy</i> , 2014, 58, 7141-7150.	1.4	8
13	Addition of E138K to R263K in HIV integrase increases resistance to dolutegravir, but fails to restore activity of the HIV integrase enzyme and viral replication capacity. <i>Journal of Antimicrobial Chemotherapy</i> , 2014, 69, 2733-2740.	1.3	47