Raymond F Schinazi

List of Publications by Year in descending order

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		22153	31849
319	13,533	59	101
papers	citations	h-index	g-index
336	336	336	14401
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Randomized Trial of Ruxolitinib in Antiretroviral-Treated Adults With Human Immunodeficiency Virus. Clinical Infectious Diseases, 2022, 74, 95-104.	5.8	31
2	GNS561 Exhibits Potent Antiviral Activity against SARS-CoV-2 through Autophagy Inhibition. Viruses, 2022, 14, 132.	3.3	10
3	Elimination of Aicardi–GoutiÔres syndrome protein SAMHD1 activates cellular innate immunity and suppresses SARS-CoV-2 replication. Journal of Biological Chemistry, 2022, 298, 101635.	3.4	9
4	Inactivation of SARS-CoV-2 and COVID-19 Patient Samples for Contemporary Immunology and Metabolomics Studies. ImmunoHorizons, 2022, 6, 144-155.	1.8	5
5	Assessment of the Abbott BinaxNOW SARS-CoV-2 rapid antigen test against viral variants of concern. IScience, 2022, 25, 103968.	4.1	14
6	The Mechanism of Action of Hepatitis B Virus Capsid Assembly Modulators Can Be Predicted from Binding to Early Assembly Intermediates. Journal of Medicinal Chemistry, 2022, 65, 4854-4864.	6.4	8
7	<i>In silico</i> design of a novel nucleotide antiviral agent by free energy perturbation. Chemical Biology and Drug Design, 2022, , .	3.2	0
8	Diastereoselective Synthesis of 2′-Dihalopyrimidine Ribonucleoside Inhibitors of Hepatitis C Virus Replication. ACS Omega, 2022, 7, 1452-1461.	3.5	1
9	The best backbone for HIV prevention, treatment, and elimination: Emtricitabine+tenofovir. Antiviral Therapy, 2022, 27, 135965352110675.	1.0	4
10	Identification of Botanical Viral Entry Inhibitors for SARSâ \in CoVâ \in 2. FASEB Journal, 2022, 36, .	0.5	0
11	HIV nucleoside reverse transcriptase inhibitors. European Journal of Medicinal Chemistry, 2022, 240, 114554.	5.5	19
12	Design, antihuman immunodeficiency activity and molecular docking studies of synthesized 2-aryl and 2-pyrimidinyl pyrrolidines. Molecular Diversity, 2021, 25, 2045-2052.	3.9	1
13	Use of Baricitinib in Patients With Moderate to Severe Coronavirus Disease 2019. Clinical Infectious Diseases, 2021, 72, 1247-1250.	5.8	116
14	Baricitinib treatment resolves lower-airway macrophage inflammation and neutrophil recruitment in SARS-CoV-2-infected rhesus macaques. Cell, 2021, 184, 460-475.e21.	28.9	156
15	COVID-19: Discovery, diagnostics and drug development. Journal of Hepatology, 2021, 74, 168-184.	3.7	302
16	Covidâ€19 will not "magically disappear― Why access to widespread testing is paramount. American Journal of Hematology, 2021, 96, 174-178.	4.1	5
17	Discovery and structure activity relationship of glyoxamide derivatives as anti-hepatitis B virus agents. Bioorganic and Medicinal Chemistry, 2021, 31, 115952.	3.0	9
18	Moving Fast Toward Hepatitis B Virus Elimination. Advances in Experimental Medicine and Biology, 2021. 1322. 115-138.	1.6	6

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19	Studies on the Efficacy, Potential Cardiotoxicity and Monkey Pharmacokinetics of GLP-26 as a Potent Hepatitis B Virus Capsid Assembly Modulator. Viruses, 2021, 13, 114.	3.3	13
20	Comparison of anti-SARS-CoV-2 activity and intracellular metabolism of remdesivir and its parent nucleoside. Current Research in Pharmacology and Drug Discovery, 2021, 2, 100045.	3.6	20
21	Disproportionate presence of adenosine in mitochondrial and chloroplast DNA of Chlamydomonas reinhardtii. IScience, 2021, 24, 102005.	4.1	5
22	Baicalein and Baicalin Inhibit SARS-CoV-2 RNA-Dependent-RNA Polymerase. Microorganisms, 2021, 9, 893.	3.6	80
23	β- <scp>d</scp> - <i>N</i> 4-hydroxycytidine Inhibits SARS-CoV-2 Through Lethal Mutagenesis But Is Also Mutagenic To Mammalian Cells. Journal of Infectious Diseases, 2021, 224, 415-419.	4.0	211
24	Non-alcoholic fatty liver disease is a risk factor for occurrence of hepatocellular carcinoma after sustained virologic response in chronic hepatitis C patients: A prospective four-years follow-up study. Metabolism Open, 2021, 10, 100090.	2.9	16
25	Pharmacokinetics of Ruxolitinib in HIV Suppressed Individuals on Antiretroviral Agent Therapy from the ACTG A5336 Study. Journal of Clinical Pharmacology, 2021, 61, 1555-1566.	2.0	1
26	Molnupiravir promotes SARS-CoV-2 mutagenesis via the RNA template. Journal of Biological Chemistry, 2021, 297, 100770.	3.4	200
27	Single-Amplicon Multiplex Real-Time Reverse Transcription-PCR with Tiled Probes To Detect SARS-CoV-2 <i>spike</i> Mutations Associated with Variants of Concern. Journal of Clinical Microbiology, 2021, 59, e0144621.	3.9	26
28	The Effect of JAK1/2 Inhibitors on HIV Reservoir Using Primary Lymphoid Cell Model of HIV Latency. Frontiers in Immunology, 2021, 12, 720697.	4.8	9
29	Structural and functional characterization explains loss of dNTPase activity of the cancer-specific R366C/H mutant SAMHD1 proteins. Journal of Biological Chemistry, 2021, 297, 101170.	3.4	7
30	RADx Variant Task Force Program for Assessing the Impact of Variants on SARS-CoV-2 Molecular and Antigen Tests. IEEE Open Journal of Engineering in Medicine and Biology, 2021, 2, 1-1.	2.3	6
31	Contemporary Approaches to the Discovery and Development of Broad-Spectrum Natural Product Prototypes for the Control of Coronaviruses. Journal of Natural Products, 2021, 84, 3001-3007.	3.0	6
32	Synthesis of 7-trifluoromethyl-7-deazapurine ribonucleoside analogs and their monophosphate prodrugs. Nucleosides, Nucleotides and Nucleic Acids, 2020, 39, 671-687.	1.1	2
33	Disentangling the lifespans of hepatitis C virusâ€infected cells and intracellular vRNA replicationâ€complexes during directâ€acting antiâ€viral therapy. Journal of Viral Hepatitis, 2020, 27, 261-269.	2.0	3
34	Intracellular metabolism and potential cardiotoxicity of a β-D-2'-C-methyl-2,6-diaminopurine ribonucleoside phosphoramidate that inhibits hepatitis C virus replication. Nucleosides, Nucleotides and Nucleic Acids, 2020, 39, 204-224.	1.1	3
35	Novel Hepatitis B Virus Capsid Assembly Modulator Induces Potent Antiviral Responses <i>In Vitro</i> and in Humanized Mice. Antimicrobial Agents and Chemotherapy, 2020, 64, .	3.2	28
36	Enhanced enzyme kinetics of reverse transcriptase variants cloned from animals infected with SIVmac239 lacking viral protein X. Journal of Biological Chemistry, 2020, 295, 16975-16986.	3.4	2

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37	Mechanistic cross-talk between DNA/RNA polymerase enzyme kinetics and nucleotide substrate availability in cells: Implications for polymerase inhibitor discovery. Journal of Biological Chemistry, 2020, 295, 13432-13443.	3.4	6
38	Application of Molecular Dynamics Simulations to the Design of Nucleotide Inhibitors Binding to Norovirus Polymerase. Journal of Chemical Information and Modeling, 2020, 60, 6566-6578.	5.4	4
39	Repurposing Nucleoside Analogs for Human Coronaviruses. Antimicrobial Agents and Chemotherapy, 2020, 65, .	3.2	45
40	Post-Catalytic Complexes with Emtricitabine or Stavudine and HIV-1 Reverse Transcriptase Reveal New Mechanistic Insights for Nucleotide Incorporation and Drug Resistance. Molecules, 2020, 25, 4868.	3.8	3
41	Response to Correspondence: Baricitinib as Treatment of COVID-19 Friend or Foe of the Pancreas? Cerda-Contreras et.al. Clinical Infectious Diseases, 2020, 73, e3978-e3979.	5.8	0
42	Novel method to quantify phenotypic markers of HIV-associated neurocognitive disorder in a murine SCID model. Journal of NeuroVirology, 2020, 26, 838-845.	2.1	2
43	Response to Correspondence: Baricitinib: Impact on Coronavirus Disease 2019 (COVID-19) Coagulopathy? Jorgensen et al. Clinical Infectious Diseases, 2020, 73, e3980-e3981.	5.8	1
44	Ribonucleotide incorporation in yeast genomic DNA shows preference for cytosine and guanosine preceded by deoxyadenosine. Nature Communications, 2020, 11, 2447.	12.8	21
45	7-Deaza-7-fluoro-2′-C-methyladenosine inhibits Zika virus infection and viral-induced neuroinflammation. Antiviral Research, 2020, 180, 104855.	4.1	8
46	Synthesis of 4′-Substituted-2′-Deoxy-2′-α-Fluoro Nucleoside Analogs as Potential Antiviral Agents. Molecules, 2020, 25, 1258.	3.8	5
47	Viral protein X reduces the incorporation of mutagenic noncanonical rNTPs during lentivirus reverse transcription in macrophages. Journal of Biological Chemistry, 2020, 295, 657-666.	3.4	3
48	Potent in vitro activity of β-D-4ʹ-chloromethyl-2ʹ-deoxy-2ʹ-fluorocytidine against Nipah virus. Antiviral Research, 2020, 175, 104712.	4.1	15
49	Ribonucleotide reductase inhibitors suppress <scp>SAMHD</scp> 1 ara― <scp>CTP</scp> ase activity enhancing cytarabine efficacy. EMBO Molecular Medicine, 2020, 12, e10419.	6.9	35
50	SAMHD1 Functions and Human Diseases. Viruses, 2020, 12, 382.	3.3	51
51	Novel 1′-homo- <i>N</i> -2′-deoxy-α-nucleosides: synthesis, characterization and biological activity. RSC Advances, 2020, 10, 15815-15824.	3.6	4
52	Disparate effects of Cytotoxic Chemotherapy on the Antiviral Activity of Antiretroviral Therapy: Implications for Treatments of HIV-Infected Cancer Patients. Antiviral Therapy, 2019, 24, 177-186.	1.0	5
53	Efficient pre-catalytic conformational change of reverse transcriptases from SAMHD1 non-counteracting primate lentiviruses during dNTP incorporation. Virology, 2019, 537, 36-44.	2.4	6
54	Structural insights into the recognition of nucleoside reverse transcriptase inhibitors by HIVâ€1 reverse transcriptase: First crystal structures with reverse transcriptase and the active triphosphate forms of lamivudine and emtricitabine. Protein Science, 2019, 28, 1664-1675.	7.6	20

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55	Nucleoside Analogs with Antiviral Activity against Yellow Fever Virus. Antimicrobial Agents and Chemotherapy, 2019, 63, .	3.2	8
56	Novel influenza polymerase PB2 inhibitors for the treatment of influenza A infection. Bioorganic and Medicinal Chemistry Letters, 2019, 29, 126639.	2.2	8
57	FRI-157-Novel HBV capsid assembly modulator inhibits pregenomic RNA encapsidation by accelerating capsid assembly kinetics and disrupting core protein dephosphorylation. Journal of Hepatology, 2019, 70, e457.	3.7	3
58	Effect of induced dNTP pool imbalance on HIV-1 reverse transcription in macrophages. Retrovirology, 2019, 16, 29.	2.0	6
59	Baricitinib reverses HIV-associated neurocognitive disorders in a SCID mouse model and reservoir seeding in vitro. Journal of Neuroinflammation, 2019, 16, 182.	7.2	36
60	Discovery of a Series of 2′-α-Fluoro,2′-β-bromo-ribonucleosides and Their Phosphoramidate Prodrugs as Potent Pan-Genotypic Inhibitors of Hepatitis C Virus. Journal of Medicinal Chemistry, 2019, 62, 1859-1874.	6.4	11
61	Potential drug–drug interactions between antiretroviral therapy and treatment regimens for multi-drug resistant tuberculosis: Implications for HIV care of MDR-TB co-infected individuals. International Journal of Infectious Diseases, 2019, 83, 98-101.	3.3	20
62	Nucleoside Analogs with Selective Antiviral Activity against Dengue Fever and Japanese Encephalitis Viruses. Antimicrobial Agents and Chemotherapy, 2019, 63, .	3.2	15
63	Synthesis of 2-fluoro-substituted and 2,6-modified purine 2′,3′-dideoxy-2′,3′-difluoro-d-arabinofuranos nucleosides from d-xylose. Tetrahedron, 2019, 75, 2037-2046.	syl _{1.9}	7
64	Visualization of Positive and Negative Sense Viral RNA for Probing the Mechanism of Direct-Acting Antivirals against Hepatitis C Virus. Viruses, 2019, 11, 1039.	3.3	14
65	Structural and Antiviral Studies of the Human Norovirus GII.4 Protease. Biochemistry, 2019, 58, 900-907.	2.5	11
66	Synthesis and anti-HCV activity of β-d-2′-deoxy-2′-α-chloro-2′-β-fluoro and β-d-2′-deoxy-2′-α-brom nucleosides and their phosphoramidate prodrugs. Bioorganic and Medicinal Chemistry, 2019, 27, 664-676.	io-2′-β 3 . 0	-fluoro 9
67	Long-term Virological and Adherence Outcomes to Antiviral Treatment in a 4-year Cohort Chronic HBV Study. Antiviral Therapy, 2019, 24, 567-579.	1.0	4
68	Mobile Health Intervention to Reduce HIV Transmission: A Randomized Trial of Behaviorally Enhanced HIV Treatment as Prevention (B-TasP). Journal of Acquired Immune Deficiency Syndromes (1999), 2018, 78, 34-42.	2.1	19
69	Acute acalculous cholecystitis during zika virus infection in an immunocompromised patient. Hepatology, 2018, 67, 2051-2054.	7.3	6
70	Towards <scp>HBV</scp> curative therapies. Liver International, 2018, 38, 102-114.	3.9	63
71	Treatment of hepatitis C virus infection with directâ€acting antiviral agents: 100% cure?. Liver International, 2018, 38, 7-13.	3.9	128
72	A research agenda for curing chronic hepatitis B virus infection. Hepatology, 2018, 67, 1127-1131.	7.3	70

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73	Host SAMHD1 protein restricts endogenous reverse transcription of HIV-1 in nondividing macrophages. Retrovirology, 2018, 15, 69.	2.0	7
74	Interplay of ancestral non-primate lentiviruses with the virus-restricting SAMHD1 proteins of their hosts. Journal of Biological Chemistry, 2018, 293, 16402-16412.	3.4	16
75	HIV transmission in discordant couples in Africa in the context of antiretroviral therapy availability. Aids, 2018, 32, 1613-1623.	2.2	5
76	Simian Immunodeficiency Virus Persistence in Cellular and Anatomic Reservoirs in Antiretroviral Therapy-Suppressed Infant Rhesus Macaques. Journal of Virology, 2018, 92, .	3.4	49
77	Template-assisted synthesis of adenine-mutagenized cDNA by a retroelement protein complex. Nucleic Acids Research, 2018, 46, 9711-9725.	14.5	21
78	Synthesis and antiviral evaluation of novel peptidomimetics as norovirus protease inhibitors. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 2165-2170.	2.2	14
79	Metabolism of Nucleosides and Nucleotides Prodrugs. Current Pharmaceutical Design, 2018, 23, 6984-7002.	1.9	8
80	Expression, Purification and Characterization of a GII.4 Norovirus Protease from Minerva Virus. Infectious Disorders - Drug Targets, 2018, 18, 224-232.	0.8	1
81	Synthesis and antiviral evaluation of novel heteroarylpyrimidines analogs as HBV capsid effectors. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 904-910.	2.2	13
82	Characterization of β- <scp>d</scp> - <i>N</i> ⁴ -Hydroxycytidine as a Novel Inhibitor of Chikungunya Virus. Antimicrobial Agents and Chemotherapy, 2017, 61, .	3.2	64
83	Synthesis and antiviral evaluation of 2′,2′,3′,3′-tetrafluoro nucleoside analogs. Tetrahedron Letters, 2017, 58, 642-644.	1.4	10
84	From <scp>HCV</scp> To <scp>HBV</scp> Cure. Liver International, 2017, 37, 73-80.	3.9	26
85	Zika in the Americas, year 2: What have we learned? What gaps remain? A report from the Global Virus Network. Antiviral Research, 2017, 144, 223-246.	4.1	104
86	2′-Chloro,2′-fluoro Ribonucleotide Prodrugs with Potent Pan-genotypic Activity against Hepatitis C Virus Replication in Culture. Journal of Medicinal Chemistry, 2017, 60, 5424-5437.	6.4	23
87	Nucleotide Substrate Specificity of Anti-Hepatitis C Virus Nucleoside Analogs for Human Mitochondrial RNA Polymerase. Antimicrobial Agents and Chemotherapy, 2017, 61, .	3.2	8
88	Increased activity of unlinked Zika virus NS2B/NS3 protease compared to linked Zika virus protease. Biochemical and Biophysical Research Communications, 2017, 492, 668-673.	2.1	21
89	Pharmacokinetics and Placental Transfer of Elvitegravir, Dolutegravir, and Other Antiretrovirals during Pregnancy. Antimicrobial Agents and Chemotherapy, 2017, 61, .	3.2	30
90	Synthesis and anti-HCV activity of a series of β- d -2′-deoxy-2′-dibromo nucleosides and their corresponding phosphoramidate prodrugs. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 5296-5299.	2.2	12

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91	Synthesis of (2 <i>S</i>)-2-Chloro-2-fluororibolactone via Stereoselective Electrophilic Fluorination. Journal of Organic Chemistry, 2017, 82, 13171-13178.	3.2	7
92	A CRISPR/Cas9 approach reveals that the polymerase activity of DNA polymerase β is dispensable for HIV-1 infection in dividing and nondividing cells. Journal of Biological Chemistry, 2017, 292, 14016-14025.	3.4	14
93	Synthesis of sulfamoylbenzamide derivatives as HBV capsid assembly effector. European Journal of Medicinal Chemistry, 2017, 138, 407-421.	5.5	40
94	Anti-human immunodeficiency activity of novel 2-arylpyrrolidine analogs. Medicinal Chemistry Research, 2017, 26, 101-108.	2.4	5
95	Synthesis and antiviral evaluation of fluorinated acyclo-nucleosides and their phosphoramidates. Nucleosides, Nucleotides and Nucleic Acids, 2017, 36, 66-82.	1.1	2
96	Ruxolitinib sensitizes ovarian cancer to reduced dose Taxol, limits tumor growth and improves survival in immune competent mice. Oncotarget, 2017, 8, 94040-94053.	1.8	14
97	Substrates and Inhibitors of SAMHD1. PLoS ONE, 2017, 12, e0169052.	2.5	45
98	Novel mechanisms to inhibit HIV reservoir seeding using Jak inhibitors. PLoS Pathogens, 2017, 13, e1006740.	4.7	71
99	Jak Inhibitors Modulate Production of Replication Competent Zika Virus in Human Hofbauer, Trophoblasts, and Neuroblastoma cells. Pathogens and Immunity, 2017, 2, 199.	3.1	22
100	HIV latency reversal research and the potential effects on the central nervous system: is concern warranted?. Journal of the International AIDS Society, 2016, 19, 21008.	3.0	0
101	Biochemical Characterization of the Active Anti-Hepatitis C Virus Metabolites of 2,6-Diaminopurine Ribonucleoside Prodrug Compared to Sofosbuvir and BMS-986094. Antimicrobial Agents and Chemotherapy, 2016, 60, 4659-4669.	3.2	11
102	Zika Virus Infects Human Placental Macrophages. Cell Host and Microbe, 2016, 20, 83-90.	11.0	410
103	SAMHD1 controls cell cycle status, apoptosis and HIV-1 infection in monocytic THP-1 cells. Virology, 2016, 495, 92-100.	2.4	77
104	Toward Elimination of Hepatitis B Virus Using Novel Drugs, Approaches, and Combined Modalities. Clinics in Liver Disease, 2016, 20, 737-749.	2.1	24
105	Efficacy and safety of 3-week response-guided triple direct-acting antiviral therapy for chronic hepatitis C infection: a phase 2, open-label, proof-of-concept study. The Lancet Gastroenterology and Hepatology, 2016, 1, 97-104.	8.1	80
106	Editorial overview: Antiviral strategies. Current Opinion in Virology, 2016, 18, v-vi.	5.4	3
107	Metabolism, Biochemical Actions, and Chemical Synthesis of Anticancer Nucleosides, Nucleotides, and Base Analogs. Chemical Reviews, 2016, 116, 14379-14455.	47.7	265
108	Sonication-Assisted Synthesis of <i>(E)</i> -2-Methyl-but-2-enyl Nucleoside Phosphonate Prodrugs. ChemistrySelect, 2016, 1, 3108-3113.	1.5	8

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109	Discovery, characterization, and lead optimization of 7-azaindole non-nucleoside HIV-1 reverse transcriptase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 4101-4105.	2.2	13
110	Synthesis and Evaluation of 2,6-Modified Purine 2′- <i>C</i> -Methyl Ribonucleosides as Inhibitors of HCV Replication. ACS Medicinal Chemistry Letters, 2016, 7, 17-22.	2.8	16
111	Metabolic profiling during HIV-1 and HIV-2 infection of primary human monocyte-derived macrophages. Virology, 2016, 491, 106-114.	2.4	32
112	A new oxygen modification cyclooctaoxygen binds to nucleic acids as sodium crown complex. Biochimica Et Biophysica Acta - General Subjects, 2016, 1860, 785-794.	2.4	0
113	The Janus kinase inhibitor ruxolitinib reduces HIV replication in human macrophages and ameliorates HIV encephalitis in a murine model. Neurobiology of Disease, 2016, 92, 137-143.	4.4	60
114	Predicting Zika virus structural biology: Challenges and opportunities for intervention. Antiviral Chemistry and Chemotherapy, 2015, 24, 118-126.	0.6	58
115	Chronic liver inflammation modifies DNA methylation at the precancerous stage of murine hepatocarcinogenesis. Oncotarget, 2015, 6, 11047-11060.	1.8	21
116	Pre-steady state kinetic analysis of HIV-1 reverse transcriptase for non-canonical ribonucleoside triphosphate incorporation and DNA synthesis from ribonucleoside-containing DNA template. Antiviral Research, 2015, 115, 75-82.	4.1	4
117	Mechanistic and Kinetic Differences between Reverse Transcriptases of Vpx Coding and Non-coding Lentiviruses. Journal of Biological Chemistry, 2015, 290, 30078-30086.	3.4	26
118	Suppression of hepatitis B virus DNA accumulation in chronically infected cells using a bacterial CRISPR/Cas RNA-guided DNA endonuclease. Virology, 2015, 476, 196-205.	2.4	202
119	Towards an HBV cure: state-of-the-art and unresolved questions—report of the ANRS workshop on HBV cure. Gut, 2015, 64, 1314-1326.	12.1	234
120	Synthesis of carbocyclic nucleoside analogs with five-membered heterocyclic nucleobases. Tetrahedron Letters, 2015, 56, 3587-3590.	1.4	7
121	Design, synthesis and evaluation of novel anti-HCV molecules that deliver intracellularly three highly potent NS5A inhibitors. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 3711-3715.	2.2	2
122	Probing the structural and molecular basis of nucleotide selectivity by human mitochondrial DNA polymerase γ. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 8596-8601.	7.1	37
123	β- <scp>d</scp> -2′- <i>C</i> -Methyl-2,6-diaminopurine Ribonucleoside Phosphoramidates are Potent and Selective Inhibitors of Hepatitis C Virus (HCV) and Are Bioconverted Intracellularly to Bioactive 2,6-Diaminopurine and Guanosine 5′-Triphosphate Forms. Journal of Medicinal Chemistry, 2015, 58, 3445-3458	6.4	30
124	Ligand similarity guided receptor selection enhances docking accuracy and recall for non-nucleoside HIV reverse transcriptase inhibitors. Journal of Molecular Modeling, 2015, 21, 282.	1.8	5
125	Synthesis and antiviral evaluation of $2\hat{a}\in^2$, $3\hat{a}\in^2$ -dideoxy- $2\hat{a}\in^2$, $3\hat{a}\in^2$ -difluoro-D-arabinofuranosyl 2,6-disubstituted purine nucleosides. Heterocyclic Communications, 2015, 21, 315-327.	1.2	9
126	Role of Marine Natural Products in the Genesis of Antiviral Agents. Chemical Reviews, 2015, 115, 9655-9706.	47.7	85

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127	Differential regulatory activities of viral protein X for anti-viral efficacy of nucleos(t)ide reverse transcriptase inhibitors in monocyte-derived macrophages and activated CD4+ T cells. Virology, 2015, 485, 313-321.	2.4	5
128	Resistance to reverse transcriptase inhibitors used in the treatment and prevention of HIV-1 infection. Future Microbiology, 2015, 10, 1773-1782.	2.0	34
129	Variation of Human Immunodeficiency Virus Type-1 Reverse Transcriptase within the Simian Immunodeficiency Virus Genome of RT-SHIV. PLoS ONE, 2014, 9, e86997.	2.5	2
130	Analysis of Multiply Spliced Transcripts in Lymphoid Tissue Reservoirs of Rhesus Macaques Infected with RT-SHIV during HAART. PLoS ONE, 2014, 9, e87914.	2.5	18
131	Residual Viremia in an RT-SHIV Rhesus Macaque HAART Model Marked by the Presence of a Predominant Plasma Clone and a Lack of Viral Evolution. PLoS ONE, 2014, 9, e88258.	2.5	9
132	Kinetic variations between reverse transcriptases of viral protein X coding and noncoding lentiviruses. Retrovirology, 2014, 11, 111.	2.0	21
133	dNTP pool modulation dynamics by SAMHD1 protein in monocyte-derived macrophages. Retrovirology, 2014, 11, 63.	2.0	36
134	<scp>HCV</scp> directâ€acting antiviral agents: the best interferonâ€free combinations. Liver International, 2014, 34, 69-78.	3.9	213
135	Anti-HIV-1 screening of (2E)-3-(2-chloro-6-methyl/methoxyquinolin-3-yl)-1-(aryl)prop-2-en-1-ones. Medicinal Chemistry Research, 2014, 23, 402-407.	2.4	7
136	Cost analysis of sofosbuvir/ribavirin versus sofosbuvir/simeprevir for genotype 1 hepatitis C virus in interferon-ineligible/intolerant individuals. Hepatology, 2014, 60, 37-45.	7.3	103
137	Molecular mechanism of HIV-1 resistance to 3′-azido-2′,3′-dideoxyguanosine. Antiviral Research, 2014, 10 62-67.)] 4.1	3
138	Ruxolitinib and Tofacitinib Are Potent and Selective Inhibitors of HIV-1 Replication and Virus Reactivation <i>In Vitro</i> . Antimicrobial Agents and Chemotherapy, 2014, 58, 1977-1986.	3.2	82
139	Chutes and ladders in hepatitis C nucleoside drug development. Antiviral Research, 2014, 102, 119-147.	4.1	69
140	Synthesis and antiviral evaluation of 2-amino-6-carbamoylpurine dioxolane nucleoside derivatives and their phosphoramidates prodrugs. Bioorganic and Medicinal Chemistry, 2014, 22, 6665-6671.	3.0	5
141	Asymmetric Binding to NS5A by Daclatasvir (BMS-790052) and Analogs Suggests Two Novel Modes of HCV Inhibition. Journal of Medicinal Chemistry, 2014, 57, 10031-10043.	6.4	44
142	Enhanced Antiretroviral Therapy in Rhesus Macaques Improves RT-SHIV Viral Decay Kinetics. Antimicrobial Agents and Chemotherapy, 2014, 58, 3927-3933.	3.2	10
143	Synthesis of Nucleoside Phosphate and Phosphonate Prodrugs. Chemical Reviews, 2014, 114, 9154-9218.	47.7	440
144	Randomized, Double-Blind, Multicenter Safety and Efficacy Study of Rifalazil Compared with Azithromycin for Treatment of Uncomplicated Genital Chlamydia trachomatis Infection in Women. Antimicrobial Agents and Chemotherapy, 2014, 58, 4014-4019.	3.2	11

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145	Approaches to hepatitis C treatment and cure using NS5A inhibitors. Infection and Drug Resistance, 2014, 7, 41.	2.7	51
146	Azetidines and spiro azetidines as novel P2 units in hepatitis C virus NS3 protease inhibitors. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 6325-6330.	2.2	17
147	Treatment as prevention and cure towards global eradication of hepatitis C virus. Trends in Microbiology, 2013, 21, 625-633.	7.7	56
148	Synthesis of Cyclopentanyl Carbocyclic 5-Fluorocytosine ((â^')-5-Fluorocarbodine) Using a Facially Selective Hydrogenation Approach. Journal of Organic Chemistry, 2013, 78, 723-727.	3.2	5
149	Synthesis and evaluation of Janus type nucleosides as potential HCV NS5B polymerase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 3385-3388.	2.2	8
150	Best strategies for global <scp>HCV</scp> eradication. Liver International, 2013, 33, 68-79.	3.9	97
151	Synthesis and evaluation of non-dimeric HCV NS5A inhibitors. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 2031-2034.	2.2	21
152	Cellular Pharmacology and Potency of HIV-1 Nucleoside Analogs in Primary Human Macrophages. Antimicrobial Agents and Chemotherapy, 2013, 57, 1262-1269.	3.2	34
153	Prodrug strategies for improved efficacy of nucleoside antiviral inhibitors. Current Opinion in HIV and AIDS, 2013, 8, 556-564.	3.8	9
154	Significance of endangered and threatened plant natural products in the control of human disease. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 16832-16837.	7.1	63
155	Preclinical Characterization of GLS4, an Inhibitor of Hepatitis B Virus Core Particle Assembly. Antimicrobial Agents and Chemotherapy, 2013, 57, 5344-5354.	3.2	99
156	The Impact of Macrophage Nucleotide Pools on HIV-1 Reverse Transcription, Viral Replication, and the Development of Novel Antiviral Agents. Molecular Biology International, 2012, 2012, 1-8.	1.7	24
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