

Evelien Am Vanderlinden

List of Publications by Year in descending order

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33
papers

954
citations

393982

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36
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36
docs citations

36
times ranked

1336
citing authors

#	ARTICLE	IF	CITATIONS
1	Novel Inhibitors of Influenza Virus Fusion: Structure-Activity Relationship and Interaction with the Viral Hemagglutinin. <i>Journal of Virology</i> , 2010, 84, 4277-4288.	1.5	137
2	Emerging Antiviral Strategies to Interfere with Influenza Virus Entry. <i>Medicinal Research Reviews</i> , 2014, 34, 301-339.	5.0	91
3	Distinct Effects of T-705 (Favipiravir) and Ribavirin on Influenza Virus Replication and Viral RNA Synthesis. <i>Antimicrobial Agents and Chemotherapy</i> , 2016, 60, 6679-6691.	1.4	86
4	Antiviral therapies on the horizon for influenza. <i>Current Opinion in Pharmacology</i> , 2016, 30, 106-115.	1.7	67
5	Anti-influenza virus activity and structure-activity relationship of aglycoristocetin derivatives with cyclobutenedione carrying hydrophobic chains. <i>Antiviral Research</i> , 2009, 82, 89-94.	1.9	49
6	Synthesis and biological evaluation of pyrimidine nucleoside monophosphate prodrugs targeted against influenza virus. <i>Antiviral Research</i> , 2012, 94, 35-43.	1.9	49
7	Microwave assisted synthesis and anti-influenza virus activity of 1-adamantyl substituted N-(1-thia-4-azaspiro[4.5]decan-4-yl)carboxamide derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2012, 20, 7155-7159.	1.4	34
8	Influenza virus entry via the GM3 ganglioside-mediated platelet-derived growth factor receptor β^2 signalling pathway. <i>Journal of General Virology</i> , 2019, 100, 583-601.	1.3	34
9	Synthesis and Anti-influenza A Virus Activity of 2,2-Dialkylamantadines and Related Compounds. <i>ACS Medicinal Chemistry Letters</i> , 2012, 3, 1065-1069.	1.3	33
10	Prodrugs of the Phosphoribosylated Forms of Hydroxypyrazinecarboxamide Pseudobase T-705 and Its De-Fluoro Analogue T-1105 as Potent Influenza Virus Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 6193-6210.	2.9	32
11	Intracytoplasmic Trapping of Influenza Virus by a Lipophilic Derivative of Aglycoristocetin. <i>Journal of Virology</i> , 2012, 86, 9416-9431.	1.5	31
12	Aniline-Based Inhibitors of Influenza H1N1 Virus Acting on Hemagglutinin-Mediated Fusion. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 98-118.	2.9	31
13	Synthesis of a cluster-forming sialylthio-d-galactose fullerene conjugate and evaluation of its interaction with influenza virus hemagglutinin and neuraminidase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 2420-2423.	1.0	28
14	Broad spectrum anti-coronavirus activity of a series of anti-malaria quinoline analogues. <i>Antiviral Research</i> , 2021, 193, 105127.	1.9	27
15	Role of the viral hemagglutinin in the anti-influenza virus activity of newly synthesized polycyclic amine compounds. <i>Antiviral Research</i> , 2013, 99, 281-291.	1.9	26
16	Cell line-dependent activation and antiviral activity of T-1105, the non-fluorinated analogue of T-705 (favipiravir). <i>Antiviral Research</i> , 2019, 167, 1-5.	1.9	25
17	Semisynthetic teicoplanin derivatives as new influenza virus binding inhibitors: Synthesis and antiviral studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 3251-3254.	1.0	23
18	Synthesis and biological evaluation of lipophilic teicoplanin pseudoaglycon derivatives containing a substituted triazole function. <i>Journal of Antibiotics</i> , 2017, 70, 152-157.	1.0	21

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19	Superior inhibition of influenza virus hemagglutinin-mediated fusion by indole-substituted spirothiazolidinones. <i>Bioorganic and Medicinal Chemistry</i> , 2020, 28, 115130.	1.4	20
20	Structure-activity relationship studies of lipophilic teicoplanin pseudoaglycon derivatives as new anti-influenza virus agents. <i>European Journal of Medicinal Chemistry</i> , 2018, 157, 1017-1030.	2.6	17
21	Synthesis and Biological Evaluation of Purine 2-Fluoro-2-deoxyriboside ProTides as Anti-Influenza Virus Agents. <i>ChemMedChem</i> , 2013, 8, 415-425.	1.6	12
22	Synthesis of fluorescent ristocetin aglycon derivatives with remarkable antibacterial and antiviral activities. <i>European Journal of Medicinal Chemistry</i> , 2012, 58, 361-367.	2.6	11
23	A few atoms make the difference: Synthetic, CD, NMR and computational studies on antiviral and antibacterial activities of glycopeptide antibiotic aglycon derivatives. <i>European Journal of Medicinal Chemistry</i> , 2015, 94, 73-86.	2.6	11
24	N-benzyl 4,4-disubstituted piperidines as a potent class of influenza H1N1 virus inhibitors showing a novel mechanism of hemagglutinin fusion peptide interaction. <i>European Journal of Medicinal Chemistry</i> , 2020, 194, 112223.	2.6	11
25	Synthesis and Structure-Activity Relationship of N-(3-Oxo-1-Thia-4-Azaspiro[4.5]Decan-4-yl)Carboxamide Inhibitors of Influenza Virus Hemagglutinin Mediated Fusion. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2015, 190, 1075-1087.	0.8	10
26	<i>In Vitro</i> Characterization of the Carbohydrate-Binding Agents HHA, GNA, and UDA as Inhibitors of Influenza A and B Virus Replication. <i>Antimicrobial Agents and Chemotherapy</i> , 2021, 65, .	1.4	8
27	Synthesis and Preliminary Biologic Evaluation of 5-Substituted-(4-substituted phenyl)-1,3-Benzoxazoles as A Novel Class of Influenza Virus A Inhibitors. <i>Chemical Biology and Drug Design</i> , 2012, 79, 1018-1024.	1.5	7
28	Discovery of Dihydro-alkoxy-benzyl-oxypyrimidines as Promising Anti-Influenza Virus Agents. <i>Chemical Biology and Drug Design</i> , 2011, 78, 596-602.	1.5	6
29	Novel N-(1-thia-4-azaspiro[4.5]decan-4-yl)carboxamide derivatives as potent and selective influenza virus fusion inhibitors. <i>Archiv Der Pharmazie</i> , 2019, 352, e1900028.	2.1	5
30	Favipiravir Does Not Inhibit Chikungunya Virus Replication in Mosquito Cells and <i>Aedes aegypti</i> Mosquitoes. <i>Microorganisms</i> , 2021, 9, 944.	1.6	4
31	A broad influenza virus inhibitor acting via IMP dehydrogenase and in synergism with ribavirin. <i>Antiviral Research</i> , 2021, 196, 105208.	1.9	4
32	Synthesis and structure-activity relationship of L-methionine-coupled 1,3,4-thiadiazole derivatives with activity against influenza virus. <i>Chemical Biology and Drug Design</i> , 2022, 99, 398-415.	1.5	3
33	4,4-Disubstituted N-benzylpiperidines: A Novel Class of Fusion Inhibitors of Influenza Virus H1N1 Targeting a New Binding Site in Hemagglutinin. <i>Proceedings (mdpi)</i> , 2019, 22, .	0.2	0