Graciela Andrei

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#	Paper	IF	Citations
402	Synthesis and antiviral activity of new pyrazole and thiazole derivatives. <i>European Journal of Medicinal Chemistry</i> , 2009 , 44, 3746-53	6.8	231
401	Comparison of membrane affinity-based method with size-exclusion chromatography for isolation of exosome-like vesicles from human plasma. <i>Journal of Translational Medicine</i> , 2018 , 16, 1	8.5	208
400	Activities of various compounds against murine and primate polyomaviruses. <i>Antimicrobial Agents and Chemotherapy</i> , 1997 , 41, 587-93	5.9	199
399	HPMPC (cidofovir), PMEA (adefovir) and Related Acyclic Nucleoside Phosphonate Analogues: A Review of their Pharmacology and Clinical Potential in the Treatment of Viral Infections. <i>Antiviral Chemistry and Chemotherapy</i> , 1997 , 8, 1-23	3.5	192
398	Synthesis and antiviral activity of imidazo[1,2-a]pyridines. <i>European Journal of Medicinal Chemistry</i> , 1999 , 34, 271-274	6.8	185
397	Highly potent and selective inhibition of varicella-zoster virus by bicyclic furopyrimidine nucleosides bearing an aryl side chain. <i>Journal of Medicinal Chemistry</i> , 2000 , 43, 4993-7	8.3	173
396	Synthesis of imidazo[1,2-a]pyridines as antiviral agents. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 5108-	12 8.3	169
395	Potent and selective inhibition of varicella-zoster virus (VZV) by nucleoside analogues with an unusual bicyclic base. <i>Journal of Medicinal Chemistry</i> , 1999 , 42, 4479-84	8.3	163
394	Structure-antiviral activity relationship in the series of pyrimidine and purine N-[2-(2-phosphonomethoxy)ethyl] nucleotide analogues. 1. Derivatives substituted at the carbon atoms of the base. <i>Journal of Medicinal Chemistry</i> , 1999 , 42, 2064-86	8.3	148
393	Successful treatment of progressive mucocutaneous infection due to acyclovir- and foscarnet-resistant herpes simplex virus with (S)-1-(3-hydroxy-2-phosphonylmethoxypropyl)cytosine (HPMPC). Clinical Infectious Diseases, 1994,	11.6	139
392	18, 570-8 Synthesis of acyclo-C-nucleosides in the imidazo[1,2-a]pyridine and pyrimidine series as antiviral agents. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 2856-9	8.3	135
391	Antiadenovirus activities of several classes of nucleoside and nucleotide analogues. <i>Antimicrobial Agents and Chemotherapy</i> , 2005 , 49, 1010-6	5.9	122
390	Synthesis and antiviral activity evaluation of some new aminoadamantane derivatives. 2. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 3307-18	8.3	122
389	6-[2-(Phosphonomethoxy)alkoxy]pyrimidines with antiviral activity. <i>Journal of Medicinal Chemistry</i> , 2002 , 45, 1918-29	8.3	120
388	The novel immunosuppressive agent mycophenolate mofetil markedly potentiates the antiherpesvirus activities of acyclovir, ganciclovir, and penciclovir in vitro and in vivo. <i>Antimicrobial Agents and Chemotherapy</i> , 1998 , 42, 216-22	5.9	117
387	Synthesis, biological evaluation, and structure analysis of a series of new 1,5-anhydrohexitol nucleosides. <i>Journal of Medicinal Chemistry</i> , 1995 , 38, 826-35	8.3	115
386	Practical and efficient synthesis of pyrano[3,2-c]pyridone, pyrano[4,3-b]pyran and their hybrids with nucleoside as potential antiviral and antileishmanial agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 809-13	2.9	113

385	Antiviral activities of 5-ethynyl-1-beta-D-ribofuranosylimidazole-4- carboxamide and related compounds. <i>Antimicrobial Agents and Chemotherapy</i> , 1991 , 35, 679-84	5.9	109
384	Topical tenofovir, a microbicide effective against HIV, inhibits herpes simplex virus-2 replication. <i>Cell Host and Microbe</i> , 2011 , 10, 379-89	23.4	97
383	Antivaccinia activities of acyclic nucleoside phosphonate derivatives in epithelial cells and organotypic cultures. <i>Antimicrobial Agents and Chemotherapy</i> , 2002 , 46, 3356-61	5.9	96
382	Treatment of anogenital papillomavirus infections with an acyclic nucleoside phosphonate analogue. <i>New England Journal of Medicine</i> , 1995 , 333, 943-4	59.2	94
381	The lantibiotic peptide labyrinthopeptin A1 demonstrates broad anti-HIV and anti-HSV activity with potential for microbicidal applications. <i>PLoS ONE</i> , 2013 , 8, e64010	3.7	91
380	5-Substituted-2,4-diamino-6-[2-(phosphonomethoxy)ethoxy]pyrimidines-acyclic nucleoside phosphonate analogues with antiviral activity. <i>Journal of Medicinal Chemistry</i> , 2003 , 46, 5064-73	8.3	90
379	Molecular approaches for the treatment of hemorrhagic fever virus infections. <i>Antiviral Research</i> , 1993 , 22, 45-75	10.8	90
378	Synthesis and antiviral properties of (+/-)-5Pnoraristeromycin and related purine carbocyclic nucleosides. A new lead for anti-human cytomegalovirus agent design. <i>Journal of Medicinal Chemistry</i> , 1992 , 35, 3372-7	8.3	89
377	Inhibitory effect of selected antiviral compounds on arenavirus replication in vitro. <i>Antiviral Research</i> , 1990 , 14, 287-99	10.8	88
376	Influence of 6- or 8-substitution on the antiviral activity of 3-arylalkylthiomethylimidazo[1,2-a]pyridine against human cytomegalovirus (CMV) and varicella-zoster virus (VZV): part II. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 9536-45	3.4	84
375	Antiviral enantiomeric preference for 5Pnoraristeromycin. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 551	-8 .3	81
374	Sulphated Polymers are Potent and Selective Inhibitors of Various Enveloped Viruses, Including Herpes Simplex Virus, Cytomegalovirus, Vesicular Stomatitis Virus, Respiratory Syncytial Virus, and Toga-, Arena- and Retroviruses. <i>Antiviral Chemistry and Chemotherapy</i> , 1990 , 1, 233-240	3.5	8o
373	Activity of a Sulfated Polysaccharide Extracted from the Red Seaweed Aghardhiella Tenera against Human Immunodeficiency Virus and Other Enveloped Viruses. <i>Antiviral Chemistry and Chemotherapy</i> , 1994 , 5, 297-303	3.5	79
372	The cyclohexene ring system as a furanose mimic: synthesis and antiviral activity of both enantiomers of cyclohexenylguanine. <i>Journal of Medicinal Chemistry</i> , 2000 , 43, 736-45	8.3	78
371	Antiviral activity of triazine analogues of 1-(S)-[3-hydroxy-2-(phosphonomethoxy)propyl]cytosine (cidofovir) and related compounds. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 1069-77	8.3	76
370	Iron chelators inhibit the growth and induce the apoptosis of Kaposiß sarcoma cells and of their putative endothelial precursors. <i>Journal of Investigative Dermatology</i> , 2000 , 115, 893-900	4.3	72
369	Antiviral activity of a sulphated polysaccharide from the red seaweed Nothogenia fastigiata. <i>Biochemical Pharmacology</i> , 1994 , 47, 2187-92	6	72
368	Synthesis and antiviral activity of 3-substituted imidazo[1,2-a]pyridines <i>Bioorganic and Medicinal Chemistry Letters</i> , 1994 , 4, 1937-1940	2.9	68

367	Comparative activity of selected antiviral compounds against clinical isolates of varicella-zoster virus. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 1995 , 14, 318-29	5.3	66
366	Acyclic nucleotide analogs derived from 8-azapurines: synthesis and antiviral activity. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 4073-88	8.3	66
365	Iron as a potential co-factor in the pathogenesis of Kaposiß sarcoma?. <i>International Journal of Cancer</i> , 1998 , 78, 720-6	7.5	64
364	Comparative activity of selected antiviral compounds against clinical isolates of human cytomegalovirus. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 1991 , 10, 1026-33	5.3	64
363	Herpes simplex virus drug-resistance: new mutations and insights. <i>Current Opinion in Infectious Diseases</i> , 2013 , 26, 551-60	5.4	63
362	New neplanocin analogues. 1. Synthesis of 6Pmodified neplanocin A derivatives as broad-spectrum antiviral agents. <i>Journal of Medicinal Chemistry</i> , 1992 , 35, 324-31	8.3	63
361	Potent, selective and cell-mediated inhibition of human herpesvirus 6 at an early stage of viral replication by the non-nucleoside compound CMV423. <i>Biochemical Pharmacology</i> , 2004 , 67, 325-36	6	61
360	Cidofovir resistance in vaccinia virus is linked to diminished virulence in mice. <i>Journal of Virology</i> , 2006 , 80, 9391-401	6.6	60
359	Synthesis and biological evaluation of acyclic 3-[(2-hydroxyethoxy)methyl] analogues of antiviral furo- and pyrrolo[2,3-d]pyrimidine nucleosides. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 4690-6	8.3	60
358	Preclinical development of bicyclic nucleoside analogues as potent and selective inhibitors of varicella zoster virus. <i>Journal of Antimicrobial Chemotherapy</i> , 2007 , 60, 1316-30	5.1	59
357	Antiviral potential of a new generation of acyclic nucleoside phosphonates, the 6-[2-(phosphonomethoxy)alkoxy]-2,4-diaminopyrimidines. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2005 , 24, 331-41	1.4	59
356	Susceptibilities of several clinical varicella-zoster virus (VZV) isolates and drug-resistant VZV strains to bicyclic furano pyrimidine nucleosides. <i>Antimicrobial Agents and Chemotherapy</i> , 2005 , 49, 1081-6	5.9	58
355	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	5.9	57
354	Synthesis and antiviral activity of 2,4-diamino-5-cyano-6-[2-(phosphonomethoxy)ethoxy]pyrimidine and related compounds. <i>Bioorganic and Medicinal Chemistry</i> , 2004 , 12, 3197-202	3.4	56
353	Current pharmacological approaches to the therapy of varicella zoster virus infections: a guide to treatment. <i>Drugs</i> , 1999 , 57, 187-206	12.1	56
352	Discovery of a new family of inhibitors of human cytomegalovirus (HCMV) based upon lipophilic alkyl furano pyrimidine dideoxy nucleosides: action via a novel non-nucleosidic mechanism. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 1847-51	8.3	54
351	Antiproliferative and apoptotic effects of iron chelators on human cervical carcinoma cells. <i>Gynecologic Oncology</i> , 2002 , 85, 95-102	4.9	54
350	Synthesis, cytostatic and anti-HIV evaluations of the new unsaturated acyclic C-5 pyrimidine nucleoside analogues. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 5624-34	3.4	53

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349	Meningoradiculoneuritis due to acyclovir-resistant varicella zoster virus in an acquired immune deficiency syndrome patient. <i>Journal of Medical Virology</i> , 1994 , 42, 338-47	19.7	53
348	Resistance of herpes simplex virus type 1 against different phosphonylmethoxyalkyl derivatives of purines and pyrimidines due to specific mutations in the viral DNA polymerase gene. <i>Journal of General Virology</i> , 2000 , 81, 639-48	4.9	53
347	CADA inhibits human immunodeficiency virus and human herpesvirus 7 replication by down-modulation of the cellular CD4 receptor. <i>Virology</i> , 2002 , 302, 342-53	3.6	52
346	Synthesis and anti-HCMV activity of 1-acyl-beta-lactams and 1-acylazetidines derived from phenylalanine. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 2253-6	2.9	51
345	Synthesis and antiviral/antitumor evaluation of 2-amino- and 2-carboxamido-3-arylsulfonylthiophenes and related compounds as a new class of diarylsulfones. <i>Bioorganic and Medicinal Chemistry</i> , 2001 , 9, 1123-32	3.4	51
344	New 2-(1-adamantylcarbonyl)pyridine and 1-acetyladamantane thiosemicarbazones-thiocarbonohydrazones: cell growth inhibitory, antiviral and antimicrobial activity evaluation. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2002 , 12, 723-7	2.9	50
343	Influence of 2-substituent on the activity of imidazo[1,2-a] pyridine derivatives against human cytomegalovirus. <i>Bioorganic and Medicinal Chemistry</i> , 2002 , 10, 941-6	3.4	49
342	Cell-dependent interference of a series of new 6-aminoquinolone derivatives with viral (HIV/CMV) transactivation. <i>Journal of Antimicrobial Chemotherapy</i> , 2005 , 56, 847-55	5.1	49
341	Acyclic/carbocyclic guanosine analogues as anti-herpesvirus agents. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2001 , 20, 271-85	1.4	49
340	Antiviral Activity of low-MW Dextran Sulphate (Derived from dextran MW 1000) Compared to Dextran Sulphate Samples of Higher MW. <i>Antiviral Chemistry and Chemotherapy</i> , 1991 , 2, 171-179	3.5	49
339	Vaccinia virus-encoded ribonucleotide reductase subunits are differentially required for replication and pathogenesis. <i>PLoS Pathogens</i> , 2010 , 6, e1000984	7.6	47
338	How Viruses Contribute to the Pathogenesis of Hemophagocytic Lymphohistiocytosis. <i>Frontiers in Immunology</i> , 2017 , 8, 1102	8.4	46
337	Anti-influenza virus activity and structure-activity relationship of aglycoristocetin derivatives with cyclobutenedione carrying hydrophobic chains. <i>Antiviral Research</i> , 2009 , 82, 89-94	10.8	46
336	Ester prodrugs of cyclic 1-(S)-[3-hydroxy-2-(phosphonomethoxy)propyl]-5-azacytosine: synthesis and antiviral activity. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 5765-72	8.3	46
335	Cidofovir Activity against Poxvirus Infections. <i>Viruses</i> , 2010 , 2, 2803-30	6.2	45
334	Characterization of herpes simplex virus type 1 thymidine kinase mutants selected under a single round of high-dose brivudin. <i>Journal of Virology</i> , 2005 , 79, 5863-9	6.6	45
333	Differential antiviral activity of derivatized dextrans. <i>Biochemical Pharmacology</i> , 1995 , 50, 743-51	6	45
332	Novel antiviral C5-substituted pyrimidine acyclic nucleoside phosphonates selected as human thymidylate kinase substrates. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 222-32	8.3	44

331	Synthesis of new C5-(1-substituted-1,2,3-triazol-4 or 5-yl)-2Pdeoxyuridines and their antiviral evaluation. <i>European Journal of Medicinal Chemistry</i> , 2011 , 46, 778-86	6.8	44
330	Polyanion inhibitors of HIV and other viruses. 7. Polyanionic compounds and polyzwitterionic compounds derived from cyclodextrins as inhibitors of HIV transmission. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 4927-32	8.3	44
329	From 1-acyl-beta-lactam human cytomegalovirus protease inhibitors to 1-benzyloxycarbonylazetidines with improved antiviral activity. A straightforward approach to convert covalent to noncovalent inhibitors. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 2612-21	8.3	44
328	Broad-spectrum antiviral activity and mechanism of antiviral action of the fluoroquinolone derivative K-12. <i>Antiviral Chemistry and Chemotherapy</i> , 1998 , 9, 403-11	3.5	44
327	Comparative activity of various compounds against clinical strains of herpes simplex virus. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 1992 , 11, 143-51	5.3	44
326	Synthesis, antiviral and anticancer activity of some novel thioureas derived from N-(4-nitro-2-phenoxyphenyl)-methanesulfonamide. <i>European Journal of Medicinal Chemistry</i> , 2009 , 44, 3591-5	6.8	43
325	In vitro-selected drug-resistant varicella-zoster virus mutants in the thymidine kinase and DNA polymerase genes yield novel phenotype-genotype associations and highlight differences between antiherpesvirus drugs. <i>Journal of Virology</i> , 2012 , 86, 2641-52	6.6	43
324	5-alkynyl analogs of arabinouridine and 2Pdeoxyuridine: cytostatic activity against herpes simplex virus and varicella-zoster thymidine kinase gene-transfected cells. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 2851-7	8.3	43
323	The N-7-substituted acyclic nucleoside analog 2-amino-7-[(1,3-dihydroxy-2-propoxy)methyl]purine is a potent and selective inhibitor of herpesvirus replication. <i>Antimicrobial Agents and Chemotherapy</i> , 1994 , 38, 2710-6	5.9	43
322	Synthesis of triterpenoid triazine derivatives from allobetulone and betulonic acid with biological activities. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 3292-300	3.4	42
321	2-Chloro-3-pyridin-3-yl-5,6,7,8-tetrahydroindolizine-1-carboxamide (CMV423), a new lead compound for the treatment of human cytomegalovirus infections. <i>Antiviral Research</i> , 2002 , 55, 413-24	10.8	42
320	Activity of the Anti-Orthopoxvirus Compound ST-246 against Vaccinia, Cowpox and Camelpox Viruses in Cell Monolayers and Organotypic Raft Cultures. <i>Antiviral Therapy</i> , 2007 , 12, 1205-1216	1.6	42
319	The novel L- and D-amino acid derivatives of hydroxyurea and hydantoins: synthesis, X-ray crystal structure study, and cytostatic and antiviral activity evaluations. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 475-82	8.3	41
318	Activities of several classes of acyclic nucleoside phosphonates against camelpox virus replication in different cell culture models. <i>Antimicrobial Agents and Chemotherapy</i> , 2007 , 51, 4410-9	5.9	40
317	Synthesis and biological evaluation of 6-(alkyn-1-yl)furo[2,3-d]pyrimidin-2(3H)-one base and nucleoside derivatives. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 391-8	8.3	40
316	New neplanocin analogues. 7. Synthesis and antiviral activity of 2-halo derivatives of neplanocin A. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 3847-52	8.3	40
315	Novel Therapeutics for Epstein?Barr Virus. <i>Molecules</i> , 2019 , 24,	4.8	39
314	Varicella-zoster virus thymidine kinase gene and antiherpetic pyrimidine nucleoside analogues in a combined gene/chemotherapy treatment for cancer. <i>Gene Therapy</i> , 1997 , 4, 1107-14	4	39

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313	Inhibitory activity of S-adenosylhomocysteine hydrolase inhibitors against human cytomegalovirus replication. <i>Antiviral Research</i> , 1993 , 21, 197-216	10.8	39
312	Synthesis and antiviral activities of 8-alkynyl-, 8-alkenyl-, and 8-alkyl-2Pdeoxyadenosine analogues. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 1307-11	8.3	39
311	The large tumor antigen: a "Swiss Army knife" protein possessing the functions required for the polyomavirus life cycle. <i>Antiviral Research</i> , 2013 , 97, 122-36	10.8	38
310	Bicyclic nucleoside inhibitors of Varicella-Zoster Virus (VZV): the effect of a terminal halogen substitution in the side-chain. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2000 , 10, 1215-7	2.9	38
309	The Low-Cost Compound Lignosulfonic Acid (LA) Exhibits Broad-Spectrum Anti-HIV and Anti-HSV Activity and Has Potential for Microbicidal Applications. <i>PLoS ONE</i> , 2015 , 10, e0131219	3.7	38
308	The novel primaquine derivatives of N-alkyl, cycloalkyl or aryl urea: synthesis, cytostatic and antiviral activity evaluations. <i>European Journal of Medicinal Chemistry</i> , 2008 , 43, 1180-7	6.8	37
307	Preparation of acyclo nucleoside phosphonate analogues based on cross-metathesis. <i>Tetrahedron</i> , 2008 , 64, 3517-3526	2.4	37
306	Novel [2Ŗ5Pbis-O-(tert-butyldimethylsilyl)-beta-D-ribofuranosyl]- 3Pspiro-5�P(4�Pamino-1�P,2�Poxathiole-2�P,2" -dioxide) derivatives with anti-HIV-1 and anti-human-cytomegalovirus activity. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 1158-68	8.3	37
305	Antitumor potential of acyclic nucleoside phosphonates. <i>Nucleosides & Nucleotides</i> , 1999 , 18, 759-71		37
304	Tricyclic analogues of acyclovir and ganciclovir. Influence of substituents in the heterocyclic moiety on the antiviral activity. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 3187-90	8.3	37
303	Novel inhibitors of human CMV. Current Opinion in Investigational Drugs, 2008, 9, 132-45		37
302	Mouse Cytomegalovirus Infection in BALB/c Mice Resembles Virus-Associated Secondary Hemophagocytic Lymphohistiocytosis and Shows a Pathogenesis Distinct from Primary Hemophagocytic Lymphohistiocytosis. <i>Journal of Immunology</i> , 2016 , 196, 3124-34	5.3	36
301	An epimer of 5Pnoraristeromycin and its antiviral properties. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 1382-4	8.3	36
300	GS-9191 is a novel topical prodrug of the nucleotide analog 9-(2-phosphonylmethoxyethyl)guanine with antiproliferative activity and possible utility in the treatment of human papillomavirus lesions. <i>Antimicrobial Agents and Chemotherapy</i> , 2009 , 53, 2777-84	5.9	35
299	4-Benzyloxy-gamma-sultone derivatives: discovery of a novel family of non-nucleoside inhibitors of human cytomegalovirus and varicella zoster virus. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 1582-91	8.3	35
298	3-deaza- and 7-deaza-5Pnoraristeromycin and their antiviral properties. <i>Journal of Medicinal Chemistry</i> , 1995 , 38, 1035-8	8.3	35
297	New neplanocin analogues. 6. Synthesis and potent antiviral activity of 6Phomoneplanocin A1. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 2392-9	8.3	35
296	Fluorescent bicyclic furo pyrimidine deoxynucleoside analogs as potent and selective inhibitors of VZV and potential future drugs for the treatment of chickenpox and shingles. <i>Drugs of the Future</i> , 2000 , 25, 1151	2.3	35

295	Activities of alkoxyalkyl esters of cidofovir (CDV), cyclic CDV, and (S)-9-(3-hydroxy-2-phosphonylmethoxypropyl)adenine against orthopoxviruses in cell monolayers and in organotypic cultures. <i>Antimicrobial Agents and Chemotherapy</i> , 2006 , 50, 2525-9	5.9	34
294	KSHV targeted therapy: an update on inhibitors of viral lytic replication. <i>Viruses</i> , 2014 , 6, 4731-59	6.2	33
293	Heterogeneity and evolution of thymidine kinase and DNA polymerase mutants of herpes simplex virus type 1: implications for antiviral therapy. <i>Journal of Infectious Diseases</i> , 2013 , 207, 1295-305	7	33
292	Mechanism of antiviral drug resistance of vaccinia virus: identification of residues in the viral DNA polymerase conferring differential resistance to antipoxvirus drugs. <i>Journal of Virology</i> , 2008 , 82, 12520	<u>0</u> 434	33
291	Synthesis and antiviral and cytostatic evaluations of the new C-5 substituted pyrimidine and furo[2,3-d]pyrimidine 4P,5Pdidehydro-L-ascorbic acid derivatives. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 4105-12	8.3	33
290	Activities of acyclic nucleoside phosphonates against Orf virus in human and ovine cell monolayers and organotypic ovine raft cultures. <i>Antimicrobial Agents and Chemotherapy</i> , 2005 , 49, 4843-52	5.9	33
289	Camelpox virus. Antiviral Research, 2011 , 92, 167-86	10.8	32
288	Three-dimensional culture models for human viral diseases and antiviral drug development. <i>Antiviral Research</i> , 2006 , 71, 96-107	10.8	32
287	Organotypic epithelial raft cultures as a model for evaluating compounds against alphaherpesviruses. <i>Antimicrobial Agents and Chemotherapy</i> , 2005 , 49, 4671-80	5.9	32
286	In vitro evaluation of the anti-orf virus activity of alkoxyalkyl esters of CDV, cCDV and (S)-HPMPA. <i>Antiviral Research</i> , 2007 , 75, 52-7	10.8	31
285	In vitro selection of drug-resistant varicella-zoster virus (VZV) mutants (OKA strain): differences between acyclovir and penciclovir?. <i>Antiviral Research</i> , 2004 , 61, 181-7	10.8	31
284	Intracellular metabolism of the N7-substituted acyclic nucleoside analog 2-amino-7-(1,3-dihydroxy-2-propoxymethyl)purine, a potent inhibitor of herpesvirus replication. <i>Molecular Pharmacology</i> , 1998 , 53, 157-65	4.3	31
283	Synthesis and antiviral activity of carbocyclic oxetanocin analogues (C-OXT-A, C-OXT-G) and related compounds. II. <i>Chemical and Pharmaceutical Bulletin</i> , 1993 , 41, 516-21	1.9	31
282	Antiviral properties of new arylsulfone derivatives with activity against human betaherpesviruses. <i>Antiviral Research</i> , 2006 , 72, 60-7	10.8	30
281	In vivo antiherpesvirus activity of N-7-substituted acyclic nucleoside analog 2-amino-7-[(1,3-dihydroxy-2-propoxy)methyl]purine. <i>Antimicrobial Agents and Chemotherapy</i> , 1995 , 39, 56-60	5.9	30
2 80	Insights into the mechanism of action of cidofovir and other acyclic nucleoside phosphonates against polyoma- and papillomaviruses and non-viral induced neoplasia. <i>Antiviral Research</i> , 2015 , 114, 21-46	10.8	29
279	Advances in the treatment of varicella-zoster virus infections. Advances in Pharmacology, 2013, 67, 107-	6§ 7	29
278	The novel phosphoramidate derivatives of NSAID 3-hydroxypropylamides: synthesis, cytostatic and antiviral activity evaluations. <i>European Journal of Medicinal Chemistry</i> , 2009 , 44, 143-51	6.8	29

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277	Furano pyrimidines as novel potent and selective anti-VZV agents. <i>Antiviral Chemistry and Chemotherapy</i> , 2001 , 12, 77-89	3.5	29	
276	Antiviral activity of anti-cytomegalovirus agents (HPMPC, HPMPA) assessed by a flow cytometric method and DNA hybridization technique. <i>Antiviral Research</i> , 1991 , 16, 1-9	10.8	29	
275	Dihydropyrimidinone/1,2,3-triazole hybrid molecules: Synthesis and anti-varicella-zoster virus (VZV) evaluation. <i>European Journal of Medicinal Chemistry</i> , 2018 , 155, 772-781	6.8	28	
274	The antiherpesvirus activity of H2G [(R)-9-[4-hydroxy-2-(hydroxymethyl)butyl]guanine] is markedly enhanced by the novel immunosuppressive agent mycophenolate mofetil. <i>Antimicrobial Agents and Chemotherapy</i> , 1998 , 42, 3285-9	5.9	28	
273	Activity of different antiviral drug combinations against human cytomegalovirus replication in vitro. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 1992 , 11, 1144-55	5.3	28	
272	Activity of the anti-orthopoxvirus compound ST-246 against vaccinia, cowpox and camelpox viruses in cell monolayers and organotypic raft cultures. <i>Antiviral Therapy</i> , 2007 , 12, 1205-16	1.6	28	
271	DNA Polymerase Mutations in Drug-Resistant Herpes Simplex Virus Mutants Determine In Vivo Neurovirulence and Drug-Enzyme Interactions. <i>Antiviral Therapy</i> , 2007 , 12, 719-732	1.6	28	
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