Robert Snoeck

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

60 15,693 103 442 h-index g-index citations papers 6.02 16,989 512 5.9 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
442	Putting drug resistant epithelial herpes keratitis in the spotlight: A case series <i>American Journal of Ophthalmology Case Reports</i> , 2022 , 25, 101268	1.3	O
441	Formulation of acyclovir (core)-dexpanthenol (sheath) nanofibrous patches for the treatment of herpes labialis. <i>International Journal of Pharmaceutics</i> , 2021 , 121354	6.5	3
440	Discovery of novel furo[2,3-d]pyrimidin-2-one-1,3,4-oxadiazole hybrid derivatives as dual antiviral and anticancer agents that induce apoptosis. <i>Archiv Der Pharmazie</i> , 2021 , 354, e2100146	4.3	9
439	Extension of furopyrimidine nucleosides with 5-alkynyl substituent: Synthesis, high fluorescence, and antiviral effect in the absence of free ribose hydroxyl groups. <i>European Journal of Medicinal Chemistry</i> , 2021 , 209, 112884	6.8	3
438	Influence of 4PSubstitution on the Activity of Gemcitabine and Its ProTide Against VZV and SARS-CoV-2. <i>ACS Medicinal Chemistry Letters</i> , 2021 , 12, 88-92	4.3	10
437	Advances and Perspectives in the Management of Varicella-Zoster Virus Infections. <i>Molecules</i> , 2021 , 26,	4.8	12
436	XPO1 inhibitors represent a novel therapeutic option in Adult T-cell Leukemia, triggering p53-mediated caspase-dependent apoptosis. <i>Blood Cancer Journal</i> , 2021 , 11, 27	7	1
435	Novel N-Substituted 3-Aryl-4-(diethoxyphosphoryl)azetidin-2-ones as Antibiotic Enhancers and Antiviral Agents in Search for a Successful Treatment of Complex Infections. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	2
434	Phenoxazine nucleoside derivatives with a multiple activity against RNA and DNA viruses. <i>European Journal of Medicinal Chemistry</i> , 2021 , 220, 113467	6.8	5
433	Peptide Derivatives of the Zonulin Inhibitor Larazotide (AT1001) as Potential Anti SARS-CoV-2: Molecular Modelling, Synthesis and Bioactivity Evaluation. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	4
432	Identification of a dual acting SARS-CoV-2 proteases inhibitor through in silico design and step-by-step biological characterization. <i>European Journal of Medicinal Chemistry</i> , 2021 , 226, 113863	6.8	4
431	Design, synthesis, chemical characterization, biological evaluation, and docking study of new 1,3,4-oxadiazole homonucleoside analogs. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2020 , 39, 1088-11	0 7 4	10
430	Amidate Prodrugs of -2-Alkylated Pyrimidine Acyclic Nucleosides Display Potent Anti-Herpesvirus Activity. <i>ACS Medicinal Chemistry Letters</i> , 2020 , 11, 1410-1415	4.3	3
429	Synthesis and Antiviral Properties of 1-Substituted 3-[E(4-Oxoquinazolin-4(3H)-yl)alkyl]uracil Derivatives. <i>Acta Naturae</i> , 2020 , 12, 134-139	2.1	0
428	Viral fitness of MHV-68 viruses harboring drug resistance mutations in the protein kinase or thymidine kinase. <i>Antiviral Research</i> , 2020 , 182, 104901	10.8	1
427	Uracil-Containing Heterodimers of a New Type: Synthesis and Study of Their Anti-Viral Properties. <i>Molecules</i> , 2020 , 25,	4.8	2
426	Synthesis of New Imidazopyridine Nucleoside Derivatives Designed as Maribavir Analogues. <i>Molecules</i> , 2020 , 25,	4.8	3

425	New acetamide derivatives containing (Ep-bromophenoxyalkyl) uracil moiety and their anticytomegalovirus activity. <i>Mendeleev Communications</i> , 2020 , 30, 602-603	1.9	Ο
424	BKTyper: Free Online Tool for Polyoma BK Virus VP1 and NCCR Typing. <i>Viruses</i> , 2020 , 12,	6.2	4
423	Design, synthesis and antiviral evaluation of novel acyclic phosphonate nucleotide analogs with triazolo[4,5-]pyridine, imidazo[4,5-]pyridine2(3)-one systems. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2020 , 39, 542-591	1.4	5
422	Substituted adamantylphthalimides: Synthesis, antiviral and antiproliferative activity. <i>Archiv Der Pharmazie</i> , 2020 , 353, e2000024	4.3	4
421	CRISPR/Cas9 Editing of the Polyomavirus Tumor Antigens Inhibits Merkel Cell Carcinoma Growth In Vitro. <i>Cancers</i> , 2019 , 11,	6.6	9
420	Antitumor and antiviral activities of 4-substituted 1,2,3-triazolyl-2,3-dibenzyl-L-ascorbic acid derivatives. <i>European Journal of Medicinal Chemistry</i> , 2019 , 184, 111739	6.8	12
419	Antiviral activity spectrum of phenoxazine nucleoside derivatives. <i>Antiviral Research</i> , 2019 , 163, 117-12-	410.8	8
418	Synthesis and anti-HSV activity of tricyclic penciclovir and hydroxybutylguanine derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2019 , 27, 1023-1033	3.4	1
417	Persistent primary cytomegalovirus infection in a kidney transplant recipient: Multi-drug resistant and compartmentalized infection leading to graft loss. <i>Antiviral Research</i> , 2019 , 168, 203-209	10.8	5
416	Acyclic nucleoside phosphonates containing the amide bond: hydroxy derivatives. <i>Monatshefte Fill Chemie</i> , 2019 , 150, 733-745	1.4	0
415	Novel Therapeutics for Epstein?Barr Virus. <i>Molecules</i> , 2019 , 24,	4.8	39
414	Synthesis and Evaluations of "1,4-Triazolyl Combretacoumarins" and Desmethoxy Analogues. <i>European Journal of Organic Chemistry</i> , 2019 , 2019, 5610-5623	3.2	4
413	Synthesis of fluorinated acyclic nucleoside phosphonates with 5-azacytosine base moiety. <i>Tetrahedron</i> , 2019 , 75, 130529	2.4	3
412	Synthesis of 4?-substituted 2?-deoxy-4?-thiocytidines and its evaluation for antineoplastic and antiviral activities. <i>Tetrahedron</i> , 2019 , 75, 4542-4555	2.4	3
411	Novel Isoxazolidine and ELactam Analogues of Homonucleosides. <i>Molecules</i> , 2019 , 24,	4.8	7
410	Synthesis, Anti-Varicella-Zoster Virus and Anti-Cytomegalovirus Activity of 4,5-Disubstituted 1,2,3-(1H)-Triazoles. <i>Medicinal Chemistry</i> , 2019 , 15, 801-812	1.8	2
409	Synthesis of uracilloumarin conjugates as potential inhibitors of virus replication. <i>Mendeleev Communications</i> , 2019 , 29, 638-639	1.9	3
408	Utilization of 1,3-Dioxolanes in the Synthesis of ⊞-branched Alkyl and Aryl 9-[2-(Phosphonomethoxy)Ethyl]Purines and Study of the Influence of ⊞-branched Substitution for Potential Biological Activity. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2019 , 38, 119-156	1.4	O

407	Highly convergent synthesis and antiviral activity of (E)-but-2-enyl nucleoside phosphonoamidates. <i>European Journal of Medicinal Chemistry</i> , 2018 , 146, 678-686	6.8	9
406	Thymidine kinase and protein kinase in drug-resistant herpesviruses: Heads of a Lernaean Hydra. <i>Drug Resistance Updates</i> , 2018 , 37, 1-16	23.2	14
405	The Anti-Human Immunodeficiency Virus Drug Tenofovir, a Reverse Transcriptase Inhibitor, Also Targets the Herpes Simplex Virus DNA Polymerase. <i>Journal of Infectious Diseases</i> , 2018 , 217, 790-801	7	7
404	Amidate Prodrugs of Cyclic 9-()-[3-Hydroxy-2-(phosphonomethoxy)propyl]adenine with Potent Anti-Herpesvirus Activity. <i>ACS Medicinal Chemistry Letters</i> , 2018 , 9, 381-385	4.3	8
403	Synthesis and antiviral evaluation of cyclopentyl nucleoside phosphonates. <i>European Journal of Medicinal Chemistry</i> , 2018 , 150, 616-625	6.8	3
402	Synthesis of a 3PC-ethynyl-Ed-ribofuranose purine nucleoside library: Discovery of C7-deazapurine analogs as potent antiproliferative nucleosides. <i>European Journal of Medicinal Chemistry</i> , 2018 , 157, 248	3-267	10
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	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
399	Lymphocyte-independent pathways underlie the pathogenesis of murine cytomegalovirus-associated secondary haemophagocytic lymphohistiocytosis. <i>Clinical and Experimental Immunology</i> , 2018 , 192, 104-119	6.2	3
398	Emimycin and its nucleoside derivatives: Synthesis and antiviral activity. <i>European Journal of Medicinal Chemistry</i> , 2018 , 144, 93-103	6.8	5
397	Xanthine-based acyclic nucleoside phosphonates with potent antiviral activity against varicella-zoster virus and human cytomegalovirus. <i>Antiviral Chemistry and Chemotherapy</i> , 2018 , 26, 2040	20661	8813050
396	Investigation of tumor-tumor interactions in a double human cervical carcinoma xenograft model in nude mice. <i>Oncotarget</i> , 2018 , 9, 21978-22000	3.3	2
395	Isoxazolidine Conjugates of N3-Substituted 6-Bromoquinazolinones-Synthesis, Anti-Varizella-Zoster Virus, and Anti-Cytomegalovirus Activity. <i>Molecules</i> , 2018 , 23,	4.8	8
394	Treatment of intraurethral condylomata acuminata with surgery and cidofovir instillations in two immunocompromised patients and review of the literature. <i>Antiviral Research</i> , 2018 , 158, 238-243	10.8	5
393	Varicella-Zoster Virus ORF9p Binding to Cellular Adaptor Protein Complex 1 Is Important for Viral Infectivity. <i>Journal of Virology</i> , 2018 , 92,	6.6	10
392	Antiviral effect of the nucleoside analogue cidofovir in the context of sexual transmission of a gammaherpesvirus in mice. <i>Journal of Antimicrobial Chemotherapy</i> , 2018 , 73, 2095-2103	5.1	1
391	Expedient synthesis and biological evaluation of alkenyl acyclic nucleoside phosphonate prodrugs. <i>Bioorganic and Medicinal Chemistry</i> , 2018 , 26, 3596-3609	3.4	2
390	Phosphonoamidate prodrugs of C5-substituted pyrimidine acyclic nucleosides for antiviral therapy. <i>Antiviral Research</i> , 2017 , 143, 262-268	10.8	9

389	Synthesis of 5,5-difluoro-5-phosphono-pent-2-en-1-yl nucleosides as potential antiviral agents. <i>RSC Advances</i> , 2017 , 7, 32282-32287	3.7	2
388	Antiviral and Cytostatic Evaluation of 5-(1-Halo-2-sulfonylvinyl)- and 5-(2-Furyl)uracil Nucleosides. Archiv Der Pharmazie, 2017 , 350, 1700023	4.3	6
387	Facile functionalization at the C4 position of pyrimidine nucleosides via amide group activation with (benzotriazol-1-yloxy)tris(dimethylamino)phosphonium hexafluorophosphate (BOP) and biological evaluations of the products. <i>Organic and Biomolecular Chemistry</i> , 2017 , 15, 1130-1139	3.9	10
386	A European multi-centre External Quality Assessment (EQA) study on phenotypic and genotypic methods used for Herpes Simplex Virus (HSV) drug resistance testing. <i>Journal of Clinical Virology</i> , 2017 , 96, 89-93	14.5	8
385	Lytic viral replication and immunopathology in a cytomegalovirus-induced mouse model of secondary hemophagocytic lymphohistiocytosis. <i>Virology Journal</i> , 2017 , 14, 240	6.1	8
384	Virtual Screening of Acyclovir Derivatives as Potential Antiviral Agents: Design, Synthesis, and Biological Evaluation of New Acyclic Nucleoside ProTides. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 7876	⁸ 7896	10
383	Design, Synthesis, and the Biological Evaluation of a New Series of Acyclic 1,2,3-Triazole Nucleosides. <i>Archiv Der Pharmazie</i> , 2017 , 350, 1700166	4.3	3
382	New prodrugs of two pyrimidine acyclic nucleoside phosphonates: Synthesis and antiviral activity. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 4637-4648	3.4	18
381	Expanding the Antiviral Spectrum of 3-Fluoro-2-(phosphonomethoxy)propyl Acyclic Nucleoside Phosphonates: Diamyl Aspartate Amidate Prodrugs. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 6220-6238	8.3	15
380	Synthesis, anti-varicella-zoster virus and anti-cytomegalovirus activity of quinazoline-2,4-diones containing isoxazolidine and phosphonate substructures. <i>European Journal of Medicinal Chemistry</i> , 2017 , 126, 84-100	6.8	17
379	Synthesis and Ativiral Activity of 5-(Benzylthio)-4-carbamyl-1,2,3-triazoles Against Human Cytomegalovirus (CMV) and Varicella-zoster Virus (VZV). <i>Medicinal Chemistry</i> , 2017 , 13, 453-464	1.8	9
378	Design, Synthesis, and Molecular Docking Studies of a Conjugated Thiadiazole-Thiourea Scaffold as Antituberculosis Agents. <i>Biological and Pharmaceutical Bulletin</i> , 2016 , 39, 502-15	2.3	24
377	Sonication-Assisted Synthesis of (E)-2-Methyl-but-2-enyl Nucleoside Phosphonate Prodrugs. <i>ChemistrySelect</i> , 2016 , 1, 3108-3113	1.8	7
376	Novel isoxazolidine analogues of homonucleosides and homonucleotides. <i>Tetrahedron</i> , 2016 , 72, 8294-8,	<u>3.0</u> .8	6
375	Acyclic nucleoside phosphonates containing the amide bond. <i>Monatshefte Fil Chemie</i> , 2016 , 147, 2163-21	7.7	2
374	Synthesis, antiviral, cytotoxic and cytostatic evaluation of -(phosphonoalkyl)uracil derivatives. Monatshefte Fil Chemie, 2016 , 147, 1081-1090	1.4	2
373	Exploring the purine core of 3PC-ethynyladenosine (EAdo) in search of novel nucleoside therapeutics. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 1970-2	2.9	6
372	Synthesis and Bioactivity of Novel Trisubstituted Triazole Nucleosides. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2016 , 35, 147-60	1.4	6

371	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
370	Basic chemokine-derived glycosaminoglycan binding peptides exert antiviral properties against dengue virus serotype 2, herpes simplex virus-1 and respiratory syncytial virus. <i>Biochemical Pharmacology</i> , 2016 , 100, 73-85	6	20
369	Cidofovir is active against human papillomavirus positive and negative head and neck and cervical tumor cells by causing DNA damage as one of its working mechanisms. <i>Oncotarget</i> , 2016 , 7, 47302-4731	18.3	6
368	New Isoxazolidine-Conjugates of Quinazolinones-Synthesis, Antiviral and Cytostatic Activity. <i>Molecules</i> , 2016 , 21,	4.8	14
367	Synthesis and the Biological Activity of Phosphonylated 1,2,3-Triazolenaphthalimide Conjugates. <i>Molecules</i> , 2016 , 21,	4.8	8
366	Resistance to the nucleotide analogue cidofovir in HPV(+) cells: a multifactorial process involving UMP/CMP kinase 1. <i>Oncotarget</i> , 2016 , 7, 10386-401	3.3	5
365	Multidrug-resistant cytomegalovirus infection in a pediatric stem cell transplantation patient. <i>Antiviral Research</i> , 2016 , 132, 149-53	10.8	5
364	Identification of an indol-based derivative as potent and selective varicella zoster virus (VZV) inhibitor. <i>European Journal of Medicinal Chemistry</i> , 2016 , 124, 773-781	6.8	11
363	Distribution and effects of amino acid changes in drug-resistant \oplus and [herpesviruses DNA polymerase. <i>Nucleic Acids Research</i> , 2016 , 44, 9530-9554	20.1	23
362	Design, Synthesis, and Antiviral Activity of Novel Ribonucleosides of 1,2,3-Triazolylbenzyl-aminophosphonates. <i>Archiv Der Pharmazie</i> , 2016 , 349, 30-41	4.3	22
361	ST-246 is a key antiviral to inhibit the viral F13L phospholipase, one of the essential proteins for orthopoxvirus wrapping. <i>Journal of Antimicrobial Chemotherapy</i> , 2015 , 70, 1367-80	5.1	18
360	Design, antiviral and cytostatic properties of isoxazolidine-containing amonafide analogues. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 3135-46	3.4	20
359	Toward the discovery of dual HCMV-VZV inhibitors: Synthesis, structure activity relationship analysis, and cytotoxicity studies of long chained 2-uracil-3-yl-N-(4-phenoxyphenyl)acetamides. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 7035-44	3.4	5
358	Human Exportin-1 is a Target for Combined Therapy of HIV and AIDS Related Lymphoma. <i>EBioMedicine</i> , 2015 , 2, 1102-13	8.8	21
357	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
356	Conservation of antiviral activity and improved selectivity in PMEO-DAPym upon pyrimidine to triazine scaffold hopping. <i>Antiviral Research</i> , 2015 , 122, 64-8	10.8	1
355	Novel halogenated 3-deazapurine, 7-deazapurine and alkylated 9-deazapurine derivatives of L-ascorbic or imino-L-ascorbic acid: Synthesis, antitumour and antiviral activity evaluations. <i>European Journal of Medicinal Chemistry</i> , 2015 , 102, 288-302	6.8	10
354	Kaposiß sarcoma-associated herpesvirus: the role of lytic replication in targeted therapy. <i>Current Opinion in Infectious Diseases</i> , 2015 , 28, 611-24	5.4	12

Phosphonylated Acyclic Guanosine Analogues with the 1,2,3-Triazole Linker. Molecules, 2015, 20, 18789-487 353 Synthesis and biological evaluation of some new 1,3,4-thiadiazole and 1,2,4-triazole derivatives from L-methionine as antituberculosis and antiviral agents. Marmara Pharmaceutical Journal, 2015, 352 10 2,88-88 Phosphonylated 8-Azahypoxantines as Acyclic Nucleotide Analogs. *Phosphorus, Sulfur and Silicon* 1 2 351 and the Related Elements, 2015, 190, 2207-2221 The Low-Cost Compound Lignosulfonic Acid (LA) Exhibits Broad-Spectrum Anti-HIV and Anti-HSV 350 38 3.7 Activity and Has Potential for Microbicidal Applications. PLoS ONE, 2015, 10, e0131219 Selective Inhibitors of Nuclear Export (SINE) Compounds Suppress Both HIV Replication and AIDS 2.2 Ο 349 Related Lymphoma. Blood, 2015, 126, 2751-2751 ProTides of N-(3-(5-(2Pdeoxyuridine))prop-2-ynyl)octanamide as potential anti-tubercular and 348 23 3.4 anti-viral agents. Bioorganic and Medicinal Chemistry, 2014, 22, 2816-24 N(4)-Acyl derivatives as lipophilic prodrugs of cidofovir and its 5-azacytosine analogue, 10 347 (S)-HPMP-5-azaC: chemistry and antiviral activity. Bioorganic and Medicinal Chemistry, **2014**, 22, 2896-906 $^{3.4}$ Synthesis of triterpenoid triazine derivatives from allobetulone and betulonic acid with biological 346 3.4 42 activities. Bioorganic and Medicinal Chemistry, 2014, 22, 3292-300 Acyclovir-resistant herpes simplex encephalitis in a patient treated with anti-tumor necrosis 16 345 14.5 factor- monoclonal antibodies. *Journal of Clinical Virology*, **2014**, 59, 67-70 Synthesis of 3P,4Pdifluoro-3Pdeoxyribonucleosides and its evaluation of the biological activities: discovery of a novel type of anti-HCV agent 3P4Pdifluorocordycepin. Bioorganic and Medicinal 344 3.4 Chemistry, 2014, 22, 6174-82 Sequestration of human cytomegalovirus by human renal and mammary epithelial cells. Virology, 3.6 9 343 2014, 460-461, 55-65 Design, synthesis, antiviral and cytostatic activity of [[1H-1,2,3-triazol-1-yl](polyhydroxy)alkylphosphonates as acyclic nucleotide analogues. Bioorganic 342 27 3.4 and Medicinal Chemistry, 2014, 22, 3629-41 An Efficient Synthesis and Antiviral Activity Evaluation of 1-[4-(5-Phenyl- 4, 5 dihydro-1H-pyrazole [& (4, 5 dihydroisoxazole)]-3-yl)-phenyl]-pyrrole- 2, 5-dione Derivates. Anti-Infective Agents, 2014, 0.6 341 12, 104-111 Antiviral Drug-Resistance Typing Reveals Compartmentalization and Dynamics of Acyclovir-Resistant Herpes Simplex Virus Type-2 (HSV-2) in a Case of Neonatal Herpes. Journal of 340 4.8 the Pediatric Infectious Diseases Society, **2014**, 3, e24-7 KAY-2-41, a novel nucleoside analogue inhibitor of orthopoxviruses in vitro and in vivo. 5.9 339 4 Antimicrobial Agents and Chemotherapy, 2014, 58, 27-37 338 KSHV targeted therapy: an update on inhibitors of viral lytic replication. Viruses, 2014, 6, 4731-59 6.2 33 Antiherpesvirus activities of two novel 4Pthiothymidine derivatives, KAY-2-41 and KAH-39-149, are dependent on viral and cellular thymidine kinases. Antimicrobial Agents and Chemotherapy, 2014, 337 5.9 11 58, 4328-40 Spectrum of activity and mechanisms of resistance of various nucleoside derivatives against 336 19 5.9 gammaherpesviruses. Antimicrobial Agents and Chemotherapy, 2014, 58, 7312-23

335	Advances in the treatment of varicella-zoster virus infections. Advances in Pharmacology, 2013, 67, 107-	68 7	29
334	Methyl-2-arylidene hydrazinecarbodithioates: synthesis and biological activity. <i>Chemical Papers</i> , 2013 , 67, 650-656	1.9	9
333	Cidofovir selectivity is based on the different response of normal and cancer cells to DNA damage. <i>BMC Medical Genomics</i> , 2013 , 6, 18	3.7	15
332	Reduced tumorigenicity and pathogenicity of cervical carcinoma SiHa cells selected for resistance to cidofovir. <i>Molecular Cancer</i> , 2013 , 12, 158	42.1	8
331	N1,N3-disubstituted uracils as nonnucleoside inhibitors of HIV-1 reverse transcriptase. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1150-8	3.4	22
330	Cidofovir treatment improves the pathology caused by the growth of human papillomavirus-positive cervical carcinoma xenografts in athymic nude mice. <i>Cancer Letters</i> , 2013 , 329, 137-45	9.9	8
329	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
328	Synthesis and anti-herpetic activity of phosphoramidate ProTides. <i>ChemMedChem</i> , 2013 , 8, 985-93	3.7	9
327	Synthesis and broad spectrum antiviral evaluation of bis(POM) prodrugs of novel acyclic nucleosides. <i>European Journal of Medicinal Chemistry</i> , 2013 , 67, 398-408	6.8	19
326	Synthesis and anti-HCMV activity of 1-[Ephenoxy)alkyl]uracil derivatives and analogues thereof. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 4151-7	3.4	19
325	A multi-targeted drug candidate with dual anti-HIV and anti-HSV activity. PLoS Pathogens, 2013, 9, e100)3 /46 6	15
324	Activity and mechanism of action of HDVD, a novel pyrimidine nucleoside derivative with high levels of selectivity and potency against gammaherpesviruses. <i>Journal of Virology</i> , 2013 , 87, 3839-51	6.6	17
323	Herpes simplex virus drug-resistance: new mutations and insights. <i>Current Opinion in Infectious Diseases</i> , 2013 , 26, 551-60	5.4	63
322	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
321	Evaluation of novel acyclic nucleoside phosphonates against human and animal gammaherpesviruses revealed an altered metabolism of cyclic prodrugs upon Epstein-Barr virus reactivation in P3HR-1 cells. <i>Journal of Virology</i> , 2013 , 87, 12422-32	6.6	14
320	Emergence of cowpox: study of the virulence of clinical strains and evaluation of antivirals. <i>PLoS ONE</i> , 2013 , 8, e55808	3.7	24
319	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
318	Synthesis and antiviral evaluation of C5-substituted-(1,3-diyne)-2Pdeoxyuridines. <i>European Journal of Medicinal Chemistry</i> , 2012 , 53, 220-8	6.8	13

(2011-2012)

317	Synthesis and biological evaluation of a series of thieno-expanded tricyclic purine 2Pdeoxy nucleoside analogues. <i>Bioorganic and Medicinal Chemistry</i> , 2012 , 20, 3009-15	3.4	19
316	Synthesis and antiviral activities of hexadecyloxypropyl prodrugs of acyclic nucleoside phosphonates containing guanine or hypoxanthine and a (S)-HPMP or PEE acyclic moiety. <i>European Journal of Medicinal Chemistry</i> , 2012 , 55, 307-14	6.8	9
315	Synthesis and antiviral evaluation of bis(POM) prodrugs of (E)-[4Pphosphono-but-2Pen-1Pyl]purine nucleosides. <i>European Journal of Medicinal Chemistry</i> , 2012 , 57, 126-33	6.8	12
314	Next-generation treatment strategies for human papillomavirus-related head and neck squamous cell carcinoma: where do we go?. <i>Reviews in Medical Virology</i> , 2012 , 22, 88-105	11.7	29
313	Synthesis and biological properties of C-2 triazolylinosine derivatives. <i>Journal of Organic Chemistry</i> , 2012 , 77, 5870-83	4.2	23
312	Skin mild hypoxia enhances killing of UVB-damaged keratinocytes through reactive oxygen species-mediated apoptosis requiring Noxa and Bim. <i>Free Radical Biology and Medicine</i> , 2012 , 52, 1111-2	2 0 .8	13
311	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
310	Mutations conferring resistance to viral DNA polymerase inhibitors in camelpox virus give different drug-susceptibility profiles in vaccinia virus. <i>Journal of Virology</i> , 2012 , 86, 7310-25	6.6	13
309	Emerging drugs for varicella-zoster virus infections. Expert Opinion on Emerging Drugs, 2011, 16, 507-35	3.7	24
308	Carbocyclic 5Pnor "reverse" fleximers. Design, synthesis, and preliminary biological activity. <i>MedChemComm</i> , 2011 , 2,	5	16
307	Dipeptidyl peptidase IV dependent water-soluble prodrugs of highly lipophilic bicyclic nucleoside analogues. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 1927-42	8.3	13
306	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
305	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
304	Synthesis of O2- and N3-(2-Phosphonomethoxy)ethyl Derivatives of 6-Phenyl- and 6-Pyridinyl-5-azacytosine. <i>Heterocycles</i> , 2011 , 83, 797	0.8	3
303	Anti-HCMV Compounds. Methods and Principles in Medicinal Chemistry, 2011, 227-282	0.4	1
302	Camelpox virus. <i>Antiviral Research</i> , 2011 , 92, 167-86	10.8	32
301	Synthesis and SAR studies on azetidine-containing dipeptides as HCMV inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 1155-61	3.4	13
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45	S-Adenosyl-L-homocysteine Hydrolase Inhibitors as Anti-Viral Agents: 5?-Deoxyaristeromycin. <i>Nucleosides & Nucleotides</i> , 1993 , 12, 185-198		26
44	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
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41	Efficacy of (S)-1-(3-hydroxy-2-phosphonylmethoxypropyl)-cytosine and 9-(1,3-dihydroxy-2-propoxymethyl)-guanine in the treatment of intracerebral murine cytomegalovirus infections in immunocompetent and immunodeficient mice. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 1993 , 12, 269-79	5.3	37
40	Poly(Hydroxy)Carboxylates as Selective Inhibitors of Cytomegalovirus and Herpes Simplex Virus Replication. <i>Antiviral Chemistry and Chemotherapy</i> , 1992 , 3, 215-222	3.5	25
39	New Polyacetal Polysulphate Active against Human Immunodeficiency Virus and other Enveloped Viruses. <i>Antiviral Chemistry and Chemotherapy</i> , 1992 , 3, 351-360	3.5	18
38	Therapy for herpesvirus infections. <i>Current Opinion in Infectious Diseases</i> , 1992 , 5, 816-826	5.4	3
37	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
36	Flow cytometric method for the detection of gpI antigens of varicella zoster virus and evaluation of anti-VZV agents. <i>Journal of Virological Methods</i> , 1992 , 38, 243-54	2.6	21
35	Synthesis and antiviral activity of acyclic nucleosides with a 3(S),5-dihydroxypentyl or 4(R)-methoxy-3(S),5-dihydroxypentyl side chain. <i>Journal of Medicinal Chemistry</i> , 1992 , 35, 1458-65	8.3	44
34	(+-)-carbocyclic 5Pnor-2Pdeoxyguanosine and related purine derivatives: synthesis and antiviral properties. <i>Journal of Medicinal Chemistry</i> , 1992 , 35, 2191-5	8.3	28
33	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
32	Synthesis and anti-herpes activity of 5-trifluorovinyl-2?-deoxyuridine. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1992 , 2, 1057-1062	2.9	6
31	Sulfated polymers inhibit the interaction of human cytomegalovirus with cell surface heparan sulfate. <i>Virology</i> , 1992 , 189, 48-58	3.6	148
30	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

29	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	
28	9-[(2RS)-3-fluoro-2-phosphonylmethoxypropyl] derivatives of purines: a class of highly selective antiretroviral agents in vitro and in vivo. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1991 , 88, 4961-5	11.5	83	
27	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	
26	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	
25	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	
24	Sulphated Cyclodextrins are Potent anti-HIV Agents Acting Synergistically with 2?,3?-dideoxynucleoside Analogues. <i>Antiviral Chemistry and Chemotherapy</i> , 1991 , 2, 45-53	3.5	41	
23	Inhibitory Effects of Polycations on the Replication of Enveloped Viruses (HIV, HSV, CMV, RSV, Influenza A Virus and Togaviruses) in vitro. <i>Antiviral Chemistry and Chemotherapy</i> , 1991 , 2, 243-248	3.5	14	
22	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	
21	Particular characteristics of the anti-human cytomegalovirus activity of (S)-1-(3-hydroxy-2-phosphonylmethoxypropyl)cytosine (HPMPC) in vitro. <i>Antiviral Research</i> , 1991 , 16, 41-52	10.8	64	
20	5-(5-Bromothien-2-yl)-2Pdeoxyuridine and 5-(5-chlorothien-2-yl)-2Pdeoxyuridine are equipotent to (E)-5-(2-bromovinyl)-2Pdeoxyuridine in the inhibition of herpes simplex virus type I replication. <i>Journal of Medicinal Chemistry</i> , 1991 , 34, 2383-9	8.3	93	
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17	Selective inhibition of human cytomegalovirus DNA synthesis by (S)-1-(3-hydroxy-2-phosphonylmethoxypropyl)cytosine [(S)-HPMPC] and 9-(1,3-dihydroxy-2-propoxymethyl)guanine (DHPG). <i>Virology</i> , 1990 , 179, 41-50	3.6	90	
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15	Novel sulfated polysaccharides: dissociation of anti-human immunodeficiency virus activity from antithrombin activity. <i>Journal of Infectious Diseases</i> , 1990 , 161, 208-13	7	64	
14	E-5-(2-Chlorovinyl)-2?-Deoxycytidine: Synthesis and Antiherpetic Activity. <i>Antiviral Chemistry and Chemotherapy</i> , 1990 , 1, 35-40	3.5	7	
13	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	80	
12	Oral Treatment of Oropharyngeal Candidiasis with Nystatin Versus Ketoconazole in Cancer Patients. <i>Drug Investigation</i> , 1990 , 2, 71-75		4	

11	Production of human monoclonal IgG antibodies reacting with cytomegalovirus (CMV). <i>Journal of Immunological Methods</i> , 1990 , 130, 209-16	2.5	9
10	Acyclic nucleotide analogues: synthesis, antiviral activity and inhibitory effects on some cellular and virus-encoded enzymes in vitro. <i>Antiviral Research</i> , 1990 , 13, 295-311	10.8	139
9	Detection of immediate early, early and late antigens of human cytomegalovirus by flow cytometry. <i>Journal of Virological Methods</i> , 1989 , 26, 247-54	2.6	28
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7	Rapid and automated tetrazolium-based colorimetric assay for the detection of anti-HIV compounds. <i>Journal of Virological Methods</i> , 1988 , 20, 309-21	2.6	1498
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