

Graciela Andrei

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/8430471/graciela-andrei-publications-by-year.pdf>

Version: 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

402
papers

11,285
citations

53
h-index

81
g-index

472
ext. papers

12,441
ext. citations

5.4
avg. IF

5.89
L-index

| # | Paper | IF | Citations |
|-----|--|------|-----------|
| 402 | 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 | 0 |
| 401 | 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 |
| 400 | 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 |
| 399 | 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 |
| 398 | 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 |
| 397 | Advances and Perspectives in the Management of Varicella-Zoster Virus Infections. <i>Molecules</i> , 2021 , 26, | 4.8 | 12 |
| 396 | 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 |
| 395 | Novel N-Substituted 3-Aryl-4-(diethoxyphosphoryl)azetid-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 |
| 394 | 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 |
| 393 | 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 |
| 392 | 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 |
| 391 | 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, 1088-1107 | 1.4 | 10 |
| 390 | 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 |
| 389 | Design, Synthesis, and Biological Evaluation of Novel C5-Modified Pyrimidine Ribofuranonucleosides as Potential Antitumor or/and Antiviral Agents. <i>Medicinal Chemistry</i> , 2020 , 16, 368-384 | 1.8 | 1 |
| 388 | Synthesis and Antiviral Properties of 1-Substituted 3-[4-Oxoquinazolin-4(3H)-yl]alkyl]uracil Derivatives. <i>Acta Naturae</i> , 2020 , 12, 134-139 | 2.1 | 0 |
| 387 | 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 |
| 386 | Uracil-Containing Heterodimers of a New Type: Synthesis and Study of Their Anti-Viral Properties. <i>Molecules</i> , 2020 , 25, | 4.8 | 2 |

| | | | |
|-----|--|------|----|
| 385 | Synthesis of New Imidazopyridine Nucleoside Derivatives Designed as Maribavir Analogues. <i>Molecules</i> , 2020 , 25, | 4.8 | 3 |
| 384 | New acetamide derivatives containing (E)-bromophenoxyalkyl)uracil moiety and their anticytomegalovirus activity. <i>Mendeleev Communications</i> , 2020 , 30, 602-603 | 1.9 | 0 |
| 383 | BKTyper: Free Online Tool for Polyoma BK Virus VP1 and NCCR Typing. <i>Viruses</i> , 2020 , 12, | 6.2 | 4 |
| 382 | Design, synthesis and antiviral evaluation of novel acyclic phosphonate nucleotide analogs with triazolo[4,5-]pyridine, imidazo[4,5-]pyridine and imidazo[4,5-]pyridin-2(3)-one systems. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2020 , 39, 542-591 | 1.4 | 5 |
| 381 | Substituted adamantylphthalimides: Synthesis, antiviral and antiproliferative activity. <i>Archiv Der Pharmazie</i> , 2020 , 353, e2000024 | 4.3 | 4 |
| 380 | CRISPR/Cas9 Editing of the Polyomavirus Tumor Antigens Inhibits Merkel Cell Carcinoma Growth In Vitro. <i>Cancers</i> , 2019 , 11, | 6.6 | 9 |
| 379 | 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 |
| 378 | Antiviral activity spectrum of phenoxazine nucleoside derivatives. <i>Antiviral Research</i> , 2019 , 163, 117-124 | 10.8 | 8 |
| 377 | Synthesis and anti-HSV activity of tricyclic penciclovir and hydroxybutylguanine derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2019 , 27, 1023-1033 | 3.4 | 1 |
| 376 | 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 |
| 375 | Acyclic nucleoside phosphonates containing the amide bond: hydroxy derivatives. <i>Monatshefte Für Chemie</i> , 2019 , 150, 733-745 | 1.4 | 0 |
| 374 | Novel Therapeutics for Epstein-Barr Virus. <i>Molecules</i> , 2019 , 24, | 4.8 | 39 |
| 373 | 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 |
| 372 | Synthesis of fluorinated acyclic nucleoside phosphonates with 5-azacytosine base moiety. <i>Tetrahedron</i> , 2019 , 75, 130529 | 2.4 | 3 |
| 371 | Meeting report: 32nd International Conference on Antiviral Research. <i>Antiviral Research</i> , 2019 , 169, 1045-1050 | 15.0 | 1 |
| 370 | 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 |
| 369 | Novel Isoxazolidine and Lactam Analogues of Homonucleosides. <i>Molecules</i> , 2019 , 24, | 4.8 | 7 |
| 368 | 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 |

| | | | |
|-----|---|------|-----|
| 367 | Synthesis of uracil- β -oumarin conjugates as potential inhibitors of virus replication. <i>Mendeleev Communications</i> , 2019 , 29, 638-639 | 1.9 | 3 |
| 366 | Utilization of 1,3-Dioxolanes in the Synthesis of β -branched Alkyl and Aryl 9-[2-(Phosphonomethoxy)Ethyl]Purines and Study of the Influence of β -branched Substitution for Potential Biological Activity. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2019 , 38, 119-156 | 1.4 | 0 |
| 365 | 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 |
| 364 | 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 |
| 363 | 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 |
| 362 | Amidate Prodrugs of Cyclic 9- β -[3-Hydroxy-2-(phosphonomethoxy)propyl]adenine with Potent Anti-Herpesvirus Activity. <i>ACS Medicinal Chemistry Letters</i> , 2018 , 9, 381-385 | 4.3 | 8 |
| 361 | Synthesis and antiviral evaluation of cyclopentyl nucleoside phosphonates. <i>European Journal of Medicinal Chemistry</i> , 2018 , 150, 616-625 | 6.8 | 3 |
| 360 | Synthesis of a 3PC-ethynyl- β -ribofuranose purine nucleoside library: Discovery of C7-deazapurine analogs as potent antiproliferative nucleosides. <i>European Journal of Medicinal Chemistry</i> , 2018 , 157, 248-267 | 6.8 | 10 |
| 359 | Antiviral Drugs for EBV. <i>Cancers</i> , 2018 , 10, | 6.6 | 26 |
| 358 | 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 |
| 357 | 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 |
| 356 | 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 |
| 355 | Emimycin and its nucleoside derivatives: Synthesis and antiviral activity. <i>European Journal of Medicinal Chemistry</i> , 2018 , 144, 93-103 | 6.8 | 5 |
| 354 | Xanthine-based acyclic nucleoside phosphonates with potent antiviral activity against varicella-zoster virus and human cytomegalovirus. <i>Antiviral Chemistry and Chemotherapy</i> , 2018 , 26, 2040206618813050 | 3.5 | 2 |
| 353 | Synthesis of a 3'-Deoxy-C-Nucleoside Phosphonate Bearing 9-Deazaadenine as Base Moiety. <i>European Journal of Organic Chemistry</i> , 2018 , 2018, 6657-6664 | 3.2 | |
| 352 | 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 |
| 351 | Isoxazolidine Conjugates of N3-Substituted 6-Bromoquinazolinones-Synthesis, Anti-Varizella-Zoster Virus, and Anti-Cytomegalovirus Activity. <i>Molecules</i> , 2018 , 23, | 4.8 | 8 |
| 350 | 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 |

| | | | |
|-----|--|------|----|
| 349 | Expedient synthesis and biological evaluation of alkenyl acyclic nucleoside phosphonate prodrugs. <i>Bioorganic and Medicinal Chemistry</i> , 2018 , 26, 3596-3609 | 3.4 | 2 |
| 348 | Phosphonoamidate prodrugs of C5-substituted pyrimidine acyclic nucleosides for antiviral therapy. <i>Antiviral Research</i> , 2017 , 143, 262-268 | 10.8 | 9 |
| 347 | 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 |
| 346 | Antiviral and Cytostatic Evaluation of 5-(1-Halo-2-sulfonylvinyl)- and 5-(2-Furyl)uracil Nucleosides. <i>Archiv Der Pharmazie</i> , 2017 , 350, 1700023 | 4.3 | 6 |
| 345 | 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 |
| 344 | 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 |
| 343 | 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 |
| 342 | 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, 7876-7896 | 8.3 | 10 |
| 341 | 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 |
| 340 | 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 |
| 339 | 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 |
| 338 | 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 |
| 337 | How Viruses Contribute to the Pathogenesis of Hemophagocytic Lymphohistiocytosis. <i>Frontiers in Immunology</i> , 2017 , 8, 1102 | 8.4 | 46 |
| 336 | 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 |
| 335 | 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 |
| 334 | 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 |
| 333 | Sonication-Assisted Synthesis of (E)-2-Methyl-but-2-enyl Nucleoside Phosphonate Prodrugs. <i>ChemistrySelect</i> , 2016 , 1, 3108-3113 | 1.8 | 7 |
| 332 | Novel isoxazolidine analogues of homonucleosides and homonucleotides. <i>Tetrahedron</i> , 2016 , 72, 8294-8308 | 3.8 | 6 |

- 331 Acyclic nucleoside phosphonates containing the amide bond. *Monatshefte Für Chemie*, **2016**, 147, 2163-2177 2
- 330 Synthesis, antiviral, cytotoxic and cytostatic evaluation of α -(phosphonoalkyl)uracil derivatives. *Monatshefte Für Chemie*, **2016**, 147, 1081-1090 1.4 2
- 329 Exploring the purine core of 3PC-ethynyladenosine (EAdo) in search of novel nucleoside therapeutics. *Bioorganic and Medicinal Chemistry Letters*, **2016**, 26, 1970-2 2.9 6
- 328 Synthesis and Bioactivity of Novel Trisubstituted Triazole Nucleosides. *Nucleosides, Nucleotides and Nucleic Acids*, **2016**, 35, 147-60 1.4 6
- 327 Mouse Cytomegalovirus Infection in BALB/c Mice Resembles Virus-Associated Secondary Hemophagocytic Lymphohistiocytosis and Shows a Pathogenesis Distinct from Primary Hemophagocytic Lymphohistiocytosis. *Journal of Immunology*, **2016**, 196, 3124-34 5.3 36
- 326 Basic chemokine-derived glycosaminoglycan binding peptides exert antiviral properties against dengue virus serotype 2, herpes simplex virus-1 and respiratory syncytial virus. *Biochemical Pharmacology*, **2016**, 100, 73-85 6 20
- 325 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. *Oncotarget*, **2016**, 7, 47302-47318 3.3 6
- 324 New Isoxazolidine-Conjugates of Quinazolinones-Synthesis, Antiviral and Cytostatic Activity. *Molecules*, **2016**, 21, 4.8 14
- 323 Synthesis and the Biological Activity of Phosphonylated 1,2,3-Triazolenaphthalimide Conjugates. *Molecules*, **2016**, 21, 4.8 8
- 322 Resistance to the nucleotide analogue cidofovir in HPV(+) cells: a multifactorial process involving UMP/CMP kinase 1. *Oncotarget*, **2016**, 7, 10386-401 3.3 5
- 321 Multidrug-resistant cytomegalovirus infection in a pediatric stem cell transplantation patient. *Antiviral Research*, **2016**, 132, 149-53 10.8 5
- 320 Identification of an indol-based derivative as potent and selective varicella zoster virus (VZV) inhibitor. *European Journal of Medicinal Chemistry*, **2016**, 124, 773-781 6.8 11
- 319 Distribution and effects of amino acid changes in drug-resistant θ and θ herpesviruses DNA polymerase. *Nucleic Acids Research*, **2016**, 44, 9530-9554 20.1 23
- 318 Design, Synthesis, and Antiviral Activity of Novel Ribonucleosides of 1,2,3-Triazolylbenzyl-aminophosphonates. *Archiv Der Pharmazie*, **2016**, 349, 30-41 4.3 22
- 317 ST-246 is a key antiviral to inhibit the viral F13L phospholipase, one of the essential proteins for orthopoxvirus wrapping. *Journal of Antimicrobial Chemotherapy*, **2015**, 70, 1367-80 5.1 18
- 316 Design, antiviral and cytostatic properties of isoxazolidine-containing amonafide analogues. *Bioorganic and Medicinal Chemistry*, **2015**, 23, 3135-46 3.4 20
- 315 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. *Bioorganic and Medicinal Chemistry*, **2015**, 23, 7035-44 3.4 5
- 314 Human Exportin-1 is a Target for Combined Therapy of HIV and AIDS Related Lymphoma. *EBioMedicine*, **2015**, 2, 1102-13 8.8 21

| | | | |
|-----|--|------|----|
| 313 | 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 |
| 312 | 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 |
| 311 | 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 |
| 310 | Norbornane-based nucleoside and nucleotide analogues locked in North conformation. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 184-91 | 3.4 | 13 |
| 309 | Kaposi's 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 |
| 308 | Phosphonylated Acyclic Guanosine Analogues with the 1,2,3-Triazole Linker. <i>Molecules</i> , 2015 , 20, 18789-807 | 4.07 | 8 |
| 307 | 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. <i>Marmara Pharmaceutical Journal</i> , 2015 , 2, 88-88 | | 10 |
| 306 | Phosphonylated 8-Azahypoxantines as Acyclic Nucleotide Analogs. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2015 , 190, 2207-2221 | 1 | 2 |
| 305 | 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 |
| 304 | Selective Inhibitors of Nuclear Export (SINE) Compounds Suppress Both HIV Replication and AIDS Related Lymphoma. <i>Blood</i> , 2015 , 126, 2751-2751 | 2.2 | 0 |
| 303 | ProTides of N-(3-(5-(2Pdeoxyuridine))prop-2-ynyl)octanamide as potential anti-tubercular and anti-viral agents. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 2816-24 | 3.4 | 23 |
| 302 | N(4)-Acyl derivatives as lipophilic prodrugs of cidofovir and its 5-azacytosine analogue, (S)-HPMP-5-azaC: chemistry and antiviral activity. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 2896-906 | 3.4 | 10 |
| 301 | 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 |
| 300 | Acyclovir-resistant herpes simplex encephalitis in a patient treated with anti-tumor necrosis factor- α monoclonal antibodies. <i>Journal of Clinical Virology</i> , 2014 , 59, 67-70 | 14.5 | 16 |
| 299 | Synthesis of 3P4Pdifluoro-3Pdeoxyribonucleosides and its evaluation of the biological activities: discovery of a novel type of anti-HCV agent 3P4Pdifluorocordycepin. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 6174-82 | 3.4 | 2 |
| 298 | Sequestration of human cytomegalovirus by human renal and mammary epithelial cells. <i>Virology</i> , 2014 , 460-461, 55-65 | 3.6 | 9 |
| 297 | Design, synthesis, antiviral and cytostatic activity of [(1H-1,2,3-triazol-1-yl)(polyhydroxy)alkylphosphonates as acyclic nucleotide analogues. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 3629-41 | 3.4 | 27 |
| 296 | 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. <i>Anti-Infective Agents</i> , 2014 , 12, 104-111 | 0.6 | |

| | | | |
|-----|--|------|----|
| 295 | Antiviral Drug-Resistance Typing Reveals Compartmentalization and Dynamics of Acyclovir-Resistant Herpes Simplex Virus Type-2 (HSV-2) in a Case of Neonatal Herpes. <i>Journal of the Pediatric Infectious Diseases Society</i> , 2014 , 3, e24-7 | 4.8 | 8 |
| 294 | KAY-2-41, a novel nucleoside analogue inhibitor of orthopoxviruses in vitro and in vivo. <i>Antimicrobial Agents and Chemotherapy</i> , 2014 , 58, 27-37 | 5.9 | 4 |
| 293 | KSHV targeted therapy: an update on inhibitors of viral lytic replication. <i>Viruses</i> , 2014 , 6, 4731-59 | 6.2 | 33 |
| 292 | Antiherpesvirus activities of two novel 4Pthiothymidine derivatives, KAY-2-41 and KAH-39-149, are dependent on viral and cellular thymidine kinases. <i>Antimicrobial Agents and Chemotherapy</i> , 2014 , 58, 4328-40 | 5.9 | 11 |
| 291 | Spectrum of activity and mechanisms of resistance of various nucleoside derivatives against gammaherpesviruses. <i>Antimicrobial Agents and Chemotherapy</i> , 2014 , 58, 7312-23 | 5.9 | 19 |
| 290 | A single vertebrate DNA virus protein disarms invertebrate immunity to RNA virus infection. <i>ELife</i> , 2014 , 3, | 8.9 | 12 |
| 289 | Advances in the treatment of varicella-zoster virus infections. <i>Advances in Pharmacology</i> , 2013 , 67, 107-687 | 6.7 | 29 |
| 288 | Methyl-2-arylidene hydrazinecarbodithioates: synthesis and biological activity. <i>Chemical Papers</i> , 2013 , 67, 650-656 | 1.9 | 9 |
| 287 | 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 |
| 286 | Reduced tumorigenicity and pathogenicity of cervical carcinoma SiHa cells selected for resistance to cidofovir. <i>Molecular Cancer</i> , 2013 , 12, 158 | 42.1 | 8 |
| 285 | 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 |
| 284 | 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 |
| 283 | 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 |
| 282 | Synthesis and anti-herpetic activity of phosphoramidate ProTides. <i>ChemMedChem</i> , 2013 , 8, 985-93 | 3.7 | 9 |
| 281 | 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 |
| 280 | Synthesis and anti-HCMV activity of 1-[E(phenoxy)alkyl]uracil derivatives and analogues thereof. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 4151-7 | 3.4 | 19 |
| 279 | A multi-targeted drug candidate with dual anti-HIV and anti-HSV activity. <i>PLoS Pathogens</i> , 2013 , 9, e1003466 | 4.6 | 15 |
| 278 | 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 |

| | | | |
|-----|--|------|----|
| 277 | Herpes simplex virus drug-resistance: new mutations and insights. <i>Current Opinion in Infectious Diseases</i> , 2013 , 26, 551-60 | 5.4 | 63 |
| 276 | 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 |
| 275 | 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 |
| 274 | Emergence of cowpox: study of the virulence of clinical strains and evaluation of antivirals. <i>PLoS ONE</i> , 2013 , 8, e55808 | 3.7 | 24 |
| 273 | 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 |
| 272 | 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 |
| 271 | 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 |
| 270 | 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 |
| 269 | 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 |
| 268 | Synthesis and biological properties of C-2 triazolylinosine derivatives. <i>Journal of Organic Chemistry</i> , 2012 , 77, 5870-83 | 4.2 | 23 |
| 267 | 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, 1111-20 | 7.8 | 13 |
| 266 | 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 |
| 265 | 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 |
| 264 | Emerging drugs for varicella-zoster virus infections. <i>Expert Opinion on Emerging Drugs</i> , 2011 , 16, 507-35 | 3.7 | 24 |
| 263 | Carbocyclic 5Pnor "reverse" fleximers. Design, synthesis, and preliminary biological activity. <i>MedChemComm</i> , 2011 , 2, | 5 | 16 |
| 262 | 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 |
| 261 | 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 |
| 260 | 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 |

| | | | |
|-----|--|------|----|
| 259 | 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 |
| 258 | Anti-HCMV Compounds. <i>Methods and Principles in Medicinal Chemistry</i> , 2011 , 227-282 | 0.4 | 1 |
| 257 | Camelpox virus. <i>Antiviral Research</i> , 2011 , 92, 167-86 | 10.8 | 32 |
| 256 | Synthesis and SAR studies on azetidine-containing dipeptides as HCMV inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 1155-61 | 3.4 | 13 |
| 255 | Design, synthesis and biological evaluation of 2Pdeoxy-2P2Pdifluoro-5-halouridine phosphoramidate ProTides. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 4338-45 | 3.4 | 11 |
| 254 | Acyclic nucleoside phosphonates with a branched 2-(2-phosphonoethoxy)ethyl chain: efficient synthesis and antiviral activity. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 4445-53 | 3.4 | 10 |
| 253 | From β -amino- β -keto to unusual bicyclic pyridine and pyrazine heterocyclic systems: synthesis and cytostatic and antiviral activities. <i>ChemMedChem</i> , 2011 , 6, 686-97 | 3.7 | 19 |
| 252 | 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 |
| 251 | New prodrugs of Adefovir and Cidofovir. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 3527-39 | 3.4 | 22 |
| 250 | Activities of different classes of acyclic nucleoside phosphonates against BK virus in primary human renal cells. <i>Antimicrobial Agents and Chemotherapy</i> , 2011 , 55, 1961-7 | 5.9 | 18 |
| 249 | Study of camelpox virus pathogenesis in athymic nude mice. <i>PLoS ONE</i> , 2011 , 6, e21561 | 3.7 | 13 |
| 248 | Solvent-free synthesis of pyrimidine nucleoside-aminophosphonate hybrids and their biological activity evaluation. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2010 , 29, 616-27 | 1.4 | 17 |
| 247 | Synthesis and biological evaluation of acyclic nucleotide analogues with a furo[2,3-d]pyrimidin-2(3H)-one base. <i>Canadian Journal of Chemistry</i> , 2010 , 88, 628-638 | 0.9 | 14 |
| 246 | Cidofovir Activity against Poxvirus Infections. <i>Viruses</i> , 2010 , 2, 2803-30 | 6.2 | 45 |
| 245 | Vaccinia virus-encoded ribonucleotide reductase subunits are differentially required for replication and pathogenesis. <i>PLoS Pathogens</i> , 2010 , 6, e1000984 | 7.6 | 47 |
| 244 | Evaluation of novel phosphoramidate ProTides of the 2Pfluoro derivatives of a potent anti-varicella zoster virus bicyclic nucleoside analogue. <i>Antiviral Chemistry and Chemotherapy</i> , 2010 , 21, 15-31 | 3.5 | 5 |
| 243 | Synthesis of ester prodrugs of 9-(S)-[3-hydroxy-2-(phosphonomethoxy)propyl]-2,6-diaminopurine (HPMPDAP) as anti-poxvirus agents. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 6825-37 | 8.3 | 24 |
| 242 | Design, synthesis and preliminary antiviral screening of new N-phenylpyrazole and dihydroisoxazole derivatives. <i>Medicinal Chemistry Research</i> , 2010 , 19, 1025-1035 | 2.2 | 9 |

| | | | |
|-----|---|------|-----|
| 241 | Epithelial raft cultures for investigations of virus growth, pathogenesis and efficacy of antiviral agents. <i>Antiviral Research</i> , 2010 , 85, 431-49 | 10.8 | 27 |
| 240 | 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 |
| 239 | Synthesis and evaluation of antiviral, antitubercular and anticancer activities of some novel thioureas derived from 4-aminobenzohydrazide hydrazones. <i>Marmara Pharmaceutical Journal</i> , 2010 , 1, 13-20 | | 18 |
| 238 | Cidofovir 2010 , 2403-2428 | | 0 |
| 237 | Tecovirimat, a p37 envelope protein inhibitor for the treatment of smallpox infection. <i>IDrugs: the Investigational Drugs Journal</i> , 2010 , 13, 181-91 | | 5 |
| 236 | Inhibition of vaccinia virus replication by two small interfering RNAs targeting B1R and G7L genes and their synergistic combination with cidofovir. <i>Antimicrobial Agents and Chemotherapy</i> , 2009 , 53, 2579-88 | 5.8 | 17 |
| 235 | 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 |
| 234 | 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 |
| 233 | 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 |
| 232 | 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 |
| 231 | Synthesis and anti-VZV activity of 6-heteroaryl derivatives of tricyclic acyclovir and 9-[[cis-1,2-bis(hydroxymethyl)cycloprop-1-yl]methyl]guanine analogues. <i>European Journal of Medicinal Chemistry</i> , 2009 , 44, 3313-7 | 6.8 | 19 |
| 230 | Synthesis and antiviral activity of new pyrazole and thiazole derivatives. <i>European Journal of Medicinal Chemistry</i> , 2009 , 44, 3746-53 | 6.8 | 231 |
| 229 | Alkenyl substituted bicyclic nucleoside analogues retain nanomolar potency against Varicella Zoster Virus. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 3025-7 | 3.4 | 7 |
| 228 | 2PFluorosugar analogues of the highly potent anti-varicella-zoster virus bicyclic nucleoside analogue (BCNA) Cf 1743. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 6264-7 | 2.9 | 5 |
| 227 | 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 |
| 226 | Viral DNA Polymerase Inhibitors 2009 , 481-526 | | 15 |
| 225 | Intracellular metabolism of the new antiviral compound 1-(S)-[3-hydroxy-2-(phosphonomethoxy)propyl]-5-azacytosine. <i>Biochemical Pharmacology</i> , 2008 , 76, 997-1005 | 6 | 19 |
| 224 | Fluorescence-based antiviral assay for the evaluation of compounds against vaccinia virus, varicella zoster virus and human cytomegalovirus. <i>Journal of Virological Methods</i> , 2008 , 151, 66-73 | 2.6 | 21 |

| | | | |
|-----|---|------|----|
| 223 | 4"-Benzoylureido-TSAO derivatives as potent and selective non-nucleoside HCMV inhibitors. Structure-activity relationship and mechanism of antiviral action. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 5823-32 | 8.3 | 22 |
| 222 | 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, 12520-34 | 6.6 | 33 |
| 221 | 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 |
| 220 | 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 |
| 219 | 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 |
| 218 | Preparation of acyclo nucleoside phosphonate analogues based on cross-metathesis. <i>Tetrahedron</i> , 2008 , 64, 3517-3526 | 2.4 | 37 |
| 217 | Successful kinase bypass with new acyclovir phosphoramidate prodrugs. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 4364-7 | 2.9 | 18 |
| 216 | Novel inhibitors of human CMV. <i>Current Opinion in Investigational Drugs</i> , 2008 , 9, 132-45 | | 37 |
| 215 | Specific inhibition of orthopoxvirus replication by a small interfering RNA targeting the D5R gene. <i>Antiviral Therapy</i> , 2008 , 13, 357-68 | 1.6 | 8 |
| 214 | Specific targeting of the F13L protein by ST-246 affects orthopoxvirus production differently. <i>Antiviral Therapy</i> , 2008 , 13, 977-90 | 1.6 | 9 |
| 213 | Specific Inhibition of Orthopoxvirus Replication by a Small Interfering RNA Targeting the D5R Gene. <i>Antiviral Therapy</i> , 2008 , 13, 357-368 | 1.6 | 13 |
| 212 | Specific Targeting of the F13L Protein by St-246 Affects Orthopoxvirus Production Differently. <i>Antiviral Therapy</i> , 2008 , 13, 977-990 | 1.6 | 13 |
| 211 | 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 |
| 210 | Synthesis and antiviral and cytostatic evaluations of the new C-5 substituted pyrimidine and furo[2,3-d]pyrimidine 4P5Pdidehydro-L-ascorbic acid derivatives. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 4105-12 | 8.3 | 33 |
| 209 | Antiviral activity of 3-(3,5-dimethylbenzyl)uracil derivatives against HIV-1 and HCMV. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2007 , 26, 1553-8 | 1.4 | 4 |
| 208 | Influence of 6 or 8-substitution on the antiviral activity of 3-phenethylthiomethylimidazo[1,2-a]pyridine against human cytomegalovirus (HCMV) and varicella-zoster virus (VZV). <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 7209-19 | 3.4 | 23 |
| 207 | Dual infection with polyomavirus BK and acyclovir-resistant herpes simplex virus successfully treated with cidofovir in a bone marrow transplant recipient. <i>Transplant Infectious Disease</i> , 2007 , 9, 126-37 | 3.7 | 23 |
| 206 | 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 |

| | | | |
|-----|---|------|----|
| 205 | Synthesis and antiviral activity of the carbocyclic analogue of the highly potent and selective anti-VZV bicyclo furano pyrimidines. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 6485-92 | 8.3 | 21 |
| 204 | Alkyne-azide click chemistry mediated carbanucleosides synthesis. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2007 , 26, 1391-4 | 1.4 | 17 |
| 203 | Novel non-nucleoside human cytomegalovirus inhibitors based upon TSAO nucleoside derivatives: structure-activity relationships. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2007 , 26, 625-8 | 1.4 | 4 |
| 202 | Cross-metathesis mediated synthesis of new acyclic nucleoside phosphonates. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2007 , 26, 1399-402 | 1.4 | 2 |
| 201 | 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 |
| 200 | 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 |
| 199 | Inhibitory activities of three classes of acyclic nucleoside phosphonates against murine polyomavirus and primate simian virus 40 strains. <i>Antimicrobial Agents and Chemotherapy</i> , 2007 , 51, 2268-73 | 5.9 | 27 |
| 198 | 5-alkynyl analogs of arabinouridine and 2-Deoxyuridine: 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 |
| 197 | Synthesis and antiviral evaluation of 6-(alkyl-heteroaryl)furo[2,3-d]pyrimidin-2(3H)-one nucleosides and analogues with ethynyl, ethenyl, and ethyl spacers at C6 of the furopyrimidine core. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 3897-905 | 8.3 | 22 |
| 196 | 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 |
| 195 | Therapy of poxvirus infections 2007 , 375-395 | | 5 |
| 194 | DNA polymerase mutations in drug-resistant herpes simplex virus mutants determine in vivo neurovirulence and drug-enzyme interactions. <i>Antiviral Therapy</i> , 2007 , 12, 719-32 | 1.6 | 8 |
| 193 | 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 |
| 192 | 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 |
| 191 | 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 |
| 190 | Antiviral properties of new arylsulfone derivatives with activity against human betaherpesviruses. <i>Antiviral Research</i> , 2006 , 72, 60-7 | 10.8 | 30 |
| 189 | Three-dimensional culture models for human viral diseases and antiviral drug development. <i>Antiviral Research</i> , 2006 , 71, 96-107 | 10.8 | 32 |
| 188 | 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 |

| | | | |
|-----|--|-----|-----|
| 187 | Cidofovir resistance in vaccinia virus is linked to diminished virulence in mice. <i>Journal of Virology</i> , 2006 , 80, 9391-401 | 6.6 | 60 |
| 186 | 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 |
| 185 | Synthesis and biological evaluation of 5-(alkyn-1-yl)-1-(p-toluenesulfonyl)uracil derivatives. <i>Canadian Journal of Chemistry</i> , 2006 , 84, 580-586 | 0.9 | 15 |
| 184 | Synthesis and anti-HIV-1 and anti-HCMV activity of 1-substituted 3-(3,5-dimethylbenzyl)uracil derivatives. <i>Chemical and Pharmaceutical Bulletin</i> , 2006 , 54, 325-33 | 1.9 | 21 |
| 183 | Tricyclic etheno analogs of PMEG and PMEDAP: synthesis and biological activity. <i>Bioorganic and Medicinal Chemistry</i> , 2006 , 14, 8057-65 | 3.4 | 13 |
| 182 | Antiadenovirus activities of several classes of nucleoside and nucleotide analogues. <i>Antimicrobial Agents and Chemotherapy</i> , 2005 , 49, 1010-6 | 5.9 | 122 |
| 181 | Synthesis, X-ray crystal structure study, and cytostatic and antiviral evaluation of the novel cycloalkyl-N-aryl-hydroxamic acids. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 884-7 | 8.3 | 12 |
| 180 | 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 |
| 179 | The novel L- and D-amino acid derivatives of hydroxyurea and hydantoin: 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 |
| 178 | Novel [2P5Pbis-O-(tert-butyl dimethylsilyl)-beta-D-ribofuranosyl]-3Pspiro-5P(4Pamino-1P2Poxathiole-2P2"-dioxide) derivatives with anti-HIV-1 and anti-human-cytomegalovirus activity. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 1158-68 | 8.3 | 37 |
| 177 | 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 |
| 176 | The journey towards elucidating the anti-HCMV activity of alkylated bicyclic furano pyrimidines. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2005 , 24, 643-5 | 1.4 | 2 |
| 175 | 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 |
| 174 | A novel type ozonizer for wastewater treatment. <i>Journal of Electrostatics</i> , 2005 , 63, 831-836 | 1.7 | 13 |
| 173 | Bicyclic nucleoside inhibitors of Varicella-Zoster virus: the effect of branching in the p-alkylphenyl side chain. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 3791-6 | 2.9 | 16 |
| 172 | Ovine skin organotypic cultures applied to the ex vivo study of orf virus infection. <i>Veterinary Research Communications</i> , 2005 , 29 Suppl 2, 245-7 | 2.9 | 5 |
| 171 | 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 |
| 170 | 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 |

| | | | |
|-----|---|------|----|
| 169 | 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 |
| 168 | Synthesis and biological evaluation of N- and O-alkylated bicyclic furanopyrimidines as non-nucleosidic inhibitors of human cytomegalovirus. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2005 , 24, 639-41 | 1.4 | 1 |
| 167 | Antiviral and cytostatic evaluation of the novel 6-acyclic chain substituted thymine derivatives. <i>Antiviral Chemistry and Chemotherapy</i> , 2005 , 16, 327-38 | 3.5 | 19 |
| 166 | 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 |
| 165 | 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 |
| 164 | Bicyclic nucleoside inhibitors of varicella-zoster virus modified on the sugar moiety: 3P and 5P derivatives. <i>Antiviral Chemistry and Chemotherapy</i> , 2004 , 15, 333-41 | 3.5 | 7 |
| 163 | Non-nucleoside structures retain full anti-HCMV potency of the dideoxy furanopyrimidine family. <i>Antiviral Chemistry and Chemotherapy</i> , 2004 , 15, 329-32 | 3.5 | 1 |
| 162 | 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 |
| 161 | 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 |
| 160 | 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 |
| 159 | Bicyclic anti-VZV nucleosides: thieno analogues bearing an alkylphenyl side chain have reduced antiviral activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 2397-9 | 2.9 | 15 |
| 158 | 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 |
| 157 | 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 |
| 156 | Synthesis and antiviral evaluation of some 3Pfluoro bicyclic nucleoside analogues. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2004 , 23, 1-5 | 1.4 | 9 |
| 155 | Inactivity of the bicyclic pyrimidine nucleoside analogues against simian varicella virus (SVV) does not correlate with their substrate activity for SVV-encoded thymidine kinase. <i>Biochemical and Biophysical Research Communications</i> , 2004 , 315, 877-83 | 3.4 | 13 |
| 154 | 6-[2-phosphonomethoxy]alkoxy]-2,4-diaminopyrimidines: a new class of acyclic pyrimidine nucleoside phosphonates with antiviral activity. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2004 , 23, 1321-7 | 1.4 | 25 |
| 153 | Bicyclic nucleoside inhibitors of varicella-zoster virus: synthesis and biological evaluation of 2P3Pdideoxy-3Pfluoro and 2Pdeoxy-xylo derivatives. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2003 , 22, 935-7 | 1.4 | 6 |
| 152 | Iron withdrawal strategies fail to prevent the growth of SiHa-induced tumors in mice. <i>Gynecologic Oncology</i> , 2003 , 90, 91-5 | 4.9 | 12 |

| | | | |
|-----|--|------|----|
| 151 | Novel bicyclic furanopyrimidines with dual anti-VZV and -HCMV activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2003 , 13, 4511-3 | 2.9 | 10 |
| 150 | Bicyclic nucleoside inhibitors of varicella-zoster virus: 5Pchloro and 3Pchloro derivatives. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2003 , 22, 931-3 | 1.4 | 6 |
| 149 | 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 |
| 148 | Synthesis of unusual bicyclic nucleosides bearing an unsaturated side-chain, as potential inhibitors of varicella-zoster virus (VZV). <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2003 , 22, 817-9 | 1.4 | 2 |
| 147 | Synthesis and antiviral activities of 3-alkylthiomethylimidazo[1,2-b]pyridazine derivatives. <i>Antiviral Chemistry and Chemotherapy</i> , 2003 , 14, 177-82 | 3.5 | 4 |
| 146 | Metabolic and pharmacological characteristics of the bicyclic nucleoside analogues (BCNAs) as highly selective inhibitors of varicella-zoster virus (VZV). <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2003 , 22, 995-7 | 1.4 | 5 |
| 145 | Antiviral and immunomodulatory activity of the metal chelator ethylenediaminedisuccinic acid against cytomegalovirus in vitro and in vivo. <i>Antiviral Research</i> , 2002 , 55, 179-88 | 10.8 | 4 |
| 144 | 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 |
| 143 | Pridine Oxide Derivatives: Structure-Activity Relationship for Inhibition of Human Immunodeficiency Virus and Cytomegalovirus Replication in Cell Culture. <i>Helvetica Chimica Acta</i> , 2002 , 85, 2961-2974 | 2 | 26 |
| 142 | Desferrioxamine enhances AIDS-associated Kaposi's sarcoma tumor development in a xenograft model. <i>International Journal of Cancer</i> , 2002 , 100, 140-3 | 7.5 | 15 |
| 141 | New 2-(1-adamantylcarbonyl)pyridine and 1-acetyladamantane thiosemicarbazones-thiocarbonohydrazone: cell growth inhibitory, antiviral and antimicrobial activity evaluation. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2002 , 12, 723-7 | 2.9 | 50 |
| 140 | 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 |
| 139 | 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 |
| 138 | Antiproliferative and apoptotic effects of iron chelators on human cervical carcinoma cells. <i>Gynecologic Oncology</i> , 2002 , 85, 95-102 | 4.9 | 54 |
| 137 | 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 |
| 136 | Alkyloxyphenyl furano pyrimidines as potent and selective anti-VZV agents with enhanced water solubility. <i>Antiviral Chemistry and Chemotherapy</i> , 2002 , 13, 91-9 | 3.5 | 4 |
| 135 | Polysulfonates derived from metal thiolate complexes as inhibitors of HIV-1 and various other enveloped viruses in vitro. <i>Antiviral Chemistry and Chemotherapy</i> , 2002 , 13, 185-95 | 3.5 | 15 |
| 134 | Synthesis, and cytotoxic activity of N(ind)-alkoxy derivatives of antibiotic arcylarubin and dechloro-rebecamycin aglycon. <i>Journal of Antibiotics</i> , 2002 , 55, 768-73 | 3.7 | 13 |

| | | | |
|-----|--|-----|-----|
| 133 | 6-[2-(Phosphonomethoxy)alkoxy]pyrimidines with antiviral activity. <i>Journal of Medicinal Chemistry</i> , 2002 , 45, 1918-29 | 8.3 | 120 |
| 132 | Bicyclic nucleoside inhibitors of Varicella-Zoster Virus (VZV): the effect of terminal unsaturation in the side chain. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2001 , 11, 391-3 | 2.9 | 23 |
| 131 | Bicyclic anti-VZV nucleosides: Thieno analogues retain full antiviral activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2001 , 11, 2507-10 | 2.9 | 19 |
| 130 | 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 |
| 129 | Evaluating phenotype and genotype of drug-resistant strains in herpesviruses. <i>Molecular Biotechnology</i> , 2001 , 18, 155-67 | 3 | 4 |
| 128 | Novel aryl substituted bicyclic furo nucleosides as extremely potent and selective anti-VZV agents. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2001 , 20, 287-96 | 1.4 | 4 |
| 127 | Acyclic/carbocyclic guanosine analogues as anti-herpesvirus agents. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2001 , 20, 271-85 | 1.4 | 49 |
| 126 | Furano pyrimidines as novel potent and selective anti-VZV agents. <i>Antiviral Chemistry and Chemotherapy</i> , 2001 , 12, 77-89 | 3.5 | 29 |
| 125 | Synthesis and antiviral evaluation of phosphoramidate derivatives of (E)-5-(2-bromovinyl)-2Pdeoxyuridine. <i>Antiviral Chemistry and Chemotherapy</i> , 2001 , 12, 293-300 | 3.5 | 18 |
| 124 | Synthesis of imidazo[1,2-a]pyridine derivatives as antiviral agents. <i>Arzneimittelforschung</i> , 2001 , 51, 304-9 | | 3 |
| 123 | Synthesis and in vitro evaluation of novel anti-varicella-zoster virus (VZV) nucleosides. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2001 , 20, 653-6 | 1.4 | 4 |
| 122 | Bicyclic furo pyrimidine nucleosides with aryloxyphenyl and halophenyl substituted side chains as potent and selective varicella-zoster virus inhibitors. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2001 , 20, 1063-6 | 1.4 | 7 |
| 121 | Bicyclic nucleoside inhibitors of varicella-zoster virus (VZV): effect of terminal unsaturation in the side-chain. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2001 , 20, 763-6 | 1.4 | 5 |
| 120 | Synthesis and anti-HIV activity of thymidine analogues bearing a 4Pcyanovinyl group and some derivatives thereof. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2001 , 20, 1927-39 | 1.4 | 6 |
| 119 | 3-Benzamido, ureido and thioureidoimidazo[1,2-a]pyridine derivatives as potential antiviral agents. <i>Chemical and Pharmaceutical Bulletin</i> , 2001 , 49, 1631-5 | 1.9 | 11 |
| 118 | Anti-(herpes simplex virus) activity of 4?-thio-2?-deoxyuridines: a biochemical investigation for viral and cellular target enzymes. <i>Biochemical Journal</i> , 2000 , 351, 319 | 3.8 | 9 |
| 117 | Anti-(herpes simplex virus) activity of 4?-thio-2?-deoxyuridines: a biochemical investigation for viral and cellular target enzymes. <i>Biochemical Journal</i> , 2000 , 351, 319-326 | 3.8 | 17 |
| 116 | Iron chelators inhibit the growth and induce the apoptosis of Kaposi's sarcoma cells and of their putative endothelial precursors. <i>Journal of Investigative Dermatology</i> , 2000 , 115, 893-900 | 4.3 | 72 |

| | | | |
|-----|---|-----|-----|
| 115 | 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 |
| 114 | Antiviral activity of ganciclovir elaidic acid ester against herpesviruses. <i>Antiviral Research</i> , 2000 , 45, 157-67.8 | | 12 |
| 113 | Thienothiadiazine 2,2-dioxide acyclonucleosides: synthesis and antiviral activity. <i>Antiviral Chemistry and Chemotherapy</i> , 2000 , 11, 221-30 | 3.5 | 3 |
| 112 | A rapid phenotypic assay for detection of acyclovir-resistant varicella-zoster virus with mutations in the thymidine kinase open reading frame. <i>Antimicrobial Agents and Chemotherapy</i> , 2000 , 44, 873-8 | 5.9 | 18 |
| 111 | Anti-varicella-zoster virus bicyclic nucleosides: replacement of furo by pyrro base reduces antiviral potency. <i>Antiviral Chemistry and Chemotherapy</i> , 2000 , 11, 343-8 | 3.5 | 21 |
| 110 | Synthesis and anti-varicella-zoster virus activity of some novel bicyclic nucleoside inhibitors: effect of enhanced aqueous solubility. <i>Antiviral Chemistry and Chemotherapy</i> , 2000 , 11, 383-93 | 3.5 | 13 |
| 109 | Nonnucleoside human cytomegalovirus inhibitors: synthesis and antiviral evaluation of (chlorophenylmethyl)benzothiadiazine dioxide derivatives. <i>Journal of Medicinal Chemistry</i> , 2000 , 43, 3267-73 | 8.3 | 23 |
| 108 | Novel agents for the therapy of varicella-zoster virus infections. <i>Expert Opinion on Investigational Drugs</i> , 2000 , 9, 1743-51 | 5.9 | 17 |
| 107 | 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 |
| 106 | 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 |
| 105 | Monitoring drug resistance for herpesviruses. <i>Methods in Molecular Medicine</i> , 2000 , 24, 151-69 | | 5 |
| 104 | Methods in Anti-HCMV Research. <i>Methods in Molecular Medicine</i> , 2000 , 33, 129-52 | | 1 |
| 103 | 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 |
| 102 | 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 |
| 101 | Synthesis and antiviral activity of 6-benzoyl-benzoxazolin-2-one and 6-benzoyl-benzothiazolin-2-one derivatives. <i>Antiviral Chemistry and Chemotherapy</i> , 1999 , 10, 87-97 | 3.5 | 4 |
| 100 | In vitro sensitivity of Kaposi's sarcoma cells to various chemotherapeutic agents including acyclic nucleoside phosphonates. <i>Antiviral Chemistry and Chemotherapy</i> , 1999 , 10, 129-34 | 3.5 | 5 |
| 99 | New metal chelates for the photostabilisation of polyolefins. <i>Polymer Degradation and Stability</i> , 1999 , 64, 165-171 | 4.7 | 5 |
| 98 | Imidazothiadiazine dioxides: synthesis and antiviral activity. <i>Bioorganic and Medicinal Chemistry</i> , 1999 , 7, 1617-23 | 3.4 | 14 |

| | | | |
|----|---|------|-----|
| 97 | Synthesis and antiviral activity of imidazo[1,2-a]pyridines. <i>European Journal of Medicinal Chemistry</i> , 1999 , 34, 271-274 | 6.8 | 185 |
| 96 | Synthesis and antiviral and cytostatic activities of carbocyclic nucleosides incorporating a modified cyclobutane ring. Part 1: Guanosine analogues. <i>Archiv Der Pharmazie</i> , 1999 , 332, 348-52 | 4.3 | 10 |
| 95 | 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 |
| 94 | Synthesis and biological evaluation of 1,2-disubstituted carbonucleosides of 6-substituted purine and 8-azapurine. <i>Nucleosides & Nucleotides</i> , 1999 , 18, 733-4 | | 12 |
| 93 | Structure-antiviral activity relationship in the series of pyrimidine and purine N-[2-(2-phosphonmethoxy)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 |
| 92 | 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 |
| 91 | Antitumor potential of acyclic nucleoside phosphonates. <i>Nucleosides & Nucleotides</i> , 1999 , 18, 759-71 | | 37 |
| 90 | Novel potential agents for human cytomegalovirus infection: synthesis and antiviral activity evaluation of benzothiadiazine dioxide acyclonucleosides. <i>Journal of Medicinal Chemistry</i> , 1999 , 42, 1145-50 | 8.3 | 21 |
| 89 | Synthesis of imidazo[1,2-a]pyridines as antiviral agents. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 5108-13 | 8.3 | 169 |
| 88 | Superior cytostatic activity of the ganciclovir elaidic acid ester due to the prolonged intracellular retention of ganciclovir anabolites in herpes simplex virus type 1 thymidine kinase gene-transfected tumor cells. <i>Gene Therapy</i> , 1998 , 5, 419-26 | 4 | 23 |
| 87 | Iron as a potential co-factor in the pathogenesis of Kaposi's sarcoma?. <i>International Journal of Cancer</i> , 1998 , 78, 720-6 | 7.5 | 64 |
| 86 | 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 |
| 85 | Discovery of type II (covalent) inactivation of S-adenosyl-L-homocysteine hydrolase involving its "hydrolytic activity": synthesis and evaluation of dihalohomovinyl nucleoside analogues derived from adenosine. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 3078-83 | 8.3 | 21 |
| 84 | Some Properties of Blends Based on High Density Polyethylene Grafted with Di-2-Ethyl-Hexyl Fumarat. <i>Journal of Macromolecular Science - Pure and Applied Chemistry</i> , 1998 , 35, 1137-1146 | 2.2 | 1 |
| 83 | Novel carbocyclic nucleosides containing a cyclobutyl ring. Guanosine and adenosine analogues. <i>Nucleosides & Nucleotides</i> , 1998 , 17, 1237-53 | | 12 |
| 82 | Specific therapies for human papilloma virus infections. <i>Current Opinion in Infectious Diseases</i> , 1998 , 11, 733-7 | 5.4 | 13 |
| 81 | 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 |
| 80 | 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 |

| | | | |
|----|--|------|-----|
| 79 | 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 |
| 78 | 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 |
| 77 | Differential Susceptibility of Several Drug-Resistant Strains of Herpes Simplex Virus Type 2 to Various Antiviral Compounds. <i>Antiviral Chemistry and Chemotherapy</i> , 1997 , 8, 457-461 | 3.5 | 15 |
| 76 | 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 |
| 75 | Polyanion inhibitors of human immunodeficiency virus and other viruses. 6. Micelle-like anti-HIV polyanionic compounds based on a carbohydrate core. <i>Journal of Medicinal Chemistry</i> , 1997 , 40, 350-6 | 8.3 | 19 |
| 74 | New neplanocin analogues. VIII. Synthesis and biological activity of 6PC-ethyl, -ethenyl, and -ethynyl derivatives of neplanocin A. <i>Chemical and Pharmaceutical Bulletin</i> , 1997 , 45, 1163-8 | 1.9 | 9 |
| 73 | Synthesis and Biological Activity of Some 2-Aminopurine Carbonucleosides. <i>Nucleosides & Nucleotides</i> , 1997 , 16, 1337-1339 | | 5 |
| 72 | Polyanion inhibitors of human immunodeficiency virus and other viruses. 5. Telomerized anionic surfactants derived from amino acids. <i>Journal of Medicinal Chemistry</i> , 1997 , 40, 342-9 | 8.3 | 16 |
| 71 | Carbocyclic oxetanocins lacking the C-3Pmethylene. <i>Journal of Medicinal Chemistry</i> , 1997 , 40, 1401-6 | 8.3 | 7 |
| 70 | Activities of various compounds against murine and primate polyomaviruses. <i>Antimicrobial Agents and Chemotherapy</i> , 1997 , 41, 587-93 | 5.9 | 199 |
| 69 | 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 |
| 68 | Cidofovir: Induction of apoptosis in human papilloma virus (HPV)-containing cell lines. <i>Antiviral Research</i> , 1997 , 34, A72 | 10.8 | 2 |
| 67 | Novel carbocyclic nucleosides containing a cyclopentyl ring. Adenosine and uridine analogues. <i>Archiv Der Pharmazie</i> , 1997 , 330, 265-7 | 4.3 | 4 |
| 66 | Phenotypic Resistance of Herpes Simplex Virus Type 1 Strains Selected in Vitro with Antiviral Compounds and Combinations Thereof. <i>Antiviral Chemistry and Chemotherapy</i> , 1996 , 7, 270-275 | 3.5 | 5 |
| 65 | 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 |
| 64 | Polyanion inhibitors of human immunodeficiency virus and other viruses. Part 2. Polymerized anionic surfactants derived from amino acids and dipeptides. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 1626-34 | 8.3 | 18 |
| 63 | 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 |
| 62 | Acyclic nucleotide analogs derived from 8-azapurines: synthesis and antiviral activity. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 4073-88 | 8.3 | 66 |

| | | | |
|----|---|------|-----|
| 61 | Synthesis of 2Paminomethyl derivatives of N-(2-(phosphonomethoxy)ethyl) nucleotide analogues as potential antiviral agents. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 3263-8 | 8.3 | 13 |
| 60 | 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 |
| 59 | Synthesis and antiviral activity evaluation of some new aminoadamantane derivatives. 2. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 3307-18 | 8.3 | 122 |
| 58 | Synthesis and Antiviral Activity of Acyclic Nucleotide Analogues Derived from 6-(Aminomethyl)purines and Purine-6-carboxamidines. <i>Collection of Czechoslovak Chemical Communications</i> , 1996 , 61, 1525-1537 | | 21 |
| 57 | Synthesis and antiviral activity of 6-chloropurine arabinoside and its 2Pdeoxy-2Pfluoro derivative. <i>Chemical and Pharmaceutical Bulletin</i> , 1996 , 44, 2331-4 | 1.9 | 10 |
| 56 | Synthesis and Antiviral Activities of Some Novel Carbocyclic Nucleosides. <i>Nucleosides & Nucleotides</i> , 1996 , 15, 1335-1346 | | 8 |
| 55 | Patterns of resistance and sensitivity to antiviral compounds of drug-resistant strains of human cytomegalovirus selected in vitro. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 1996 , 15, 574-9 | 5.3 | 22 |
| 54 | Mechanism of the Antiviral Activity of New Aurintricarboxylic Acid Analogues. <i>Antiviral Chemistry and Chemotherapy</i> , 1996 , 7, 142-152 | 3.5 | 13 |
| 53 | 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 |
| 52 | Palladium-catalyzed synthesis of (E)-5-(2-acylviny)-2?-deoxyuridines and their antiviral and cytotoxic activities. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1995 , 5, 1627-1632 | 2.9 | 6 |
| 51 | Combination of Azidothymidine (AZT) and (E)-5-(2-Bromoviny)-2?-deoxyuridine (BVDU) Inhibits the Replication of Herpes Simplex Virus Type 1 (HSV-1) and Type 2 (HSV-2) and Varicella Zoster Virus (VZV) Strains That Are Deficient in the Expression of the Viral Thymidine Kinase (tk). <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 1995 , 14, 559-562 | 1.4 | 3 |
| 50 | Susceptibilities of several drug-resistant herpes simplex virus type 1 strains to alternative antiviral compounds. <i>Antimicrobial Agents and Chemotherapy</i> , 1995 , 39, 1632-5 | 5.9 | 25 |
| 49 | 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 |
| 48 | 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 |
| 47 | 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 |
| 46 | 3-deaza- and 7-deaza-5Pnoraristeromycin and their antiviral properties. <i>Journal of Medicinal Chemistry</i> , 1995 , 38, 1035-8 | 8.3 | 35 |
| 45 | Differential antiviral activity of derivatized dextrans. <i>Biochemical Pharmacology</i> , 1995 , 50, 743-51 | 6 | 45 |
| 44 | The cytostatic activity of 5-(1-azidoviny)-2Pdeoxyuridine (AzVDU) against herpes simplex virus thymidine kinase gene-transfected FM3A cells is due to inhibition of thymidylate synthase and enhanced by UV light ($\lambda = 254 \text{ nm}$) exposure. <i>FEBS Letters</i> , 1995 , 373, 41-4 | 3.8 | 11 |

| | | | |
|----|--|------|-----|
| 43 | Synthesis and Antiviral Activity of 2 and 3-Substituted Imidazo[1,2-a]pyrimidine. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 1995 , 14, 551-554 | 1.4 | 13 |
| 42 | Polyanion inhibitors of human immunodeficiency virus and other viruses. 1. Polymerized anionic surfactants. <i>Journal of Medicinal Chemistry</i> , 1995 , 38, 2433-40 | 8.3 | 20 |
| 41 | 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 |
| 40 | Dipyridamole Potentiates the Activity of Various Acyclic Nucleoside Phosphonates against Varicella-Zoster Virus, Herpes Simplex Virus and Human Cytomegalovirus. <i>Antiviral Chemistry and Chemotherapy</i> , 1994 , 5, 312-321 | 3.5 | 9 |
| 39 | Human Brain Tumour Cell Lines as Cell Substrate to Demonstrate Sensitivity/Resistance of Herpes Simplex Virus Types 1 and 2 to Nucleoside Analogues. <i>Antiviral Chemistry and Chemotherapy</i> , 1994 , 5, 263-270 | 3.5 | 3 |
| 38 | Activity of a Sulfated Polysaccharide Extracted from the Red Seaweed <i>Aghardhiella Tenera</i> against Human Immunodeficiency Virus and Other Enveloped Viruses. <i>Antiviral Chemistry and Chemotherapy</i> , 1994 , 5, 297-303 | 3.5 | 79 |
| 37 | Purification and Partial Characterization of an Antiviral Active Peptide from <i>Melia Azedarach</i> L.. <i>Antiviral Chemistry and Chemotherapy</i> , 1994 , 5, 105-110 | 3.5 | 10 |
| 36 | Successful treatment of progressive mucocutaneous infection due to acyclovir- and foscarnet-resistant herpes simplex virus with (S)-1-(3-hydroxy-2-phosphonylmethoxypropyl)cytosine (HPMPC). <i>Clinical Infectious Diseases</i> , 1994 , 18, 570-8 | 11.6 | 139 |
| 35 | Facile preparation of 9-H-pyrimido [4,5-b] [1,4] diazepine derivatives from 4,5-diaminopyrimidines and ethyl pyruvate.. <i>Tetrahedron</i> , 1994 , 50, 13511-13522 | 2.4 | 6 |
| 34 | 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 |
| 33 | 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 |
| 32 | 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 |
| 31 | Chemotherapy of varicella zoster virus infections. <i>International Journal of Antimicrobial Agents</i> , 1994 , 4, 211-26 | 14.3 | 16 |
| 30 | Antiviral activity of a sulphated polysaccharide from the red seaweed <i>Nothogenia fastigiata</i> . <i>Biochemical Pharmacology</i> , 1994 , 47, 2187-92 | 6 | 72 |
| 29 | An epimer of 5Pnoraristeromycin and its antiviral properties. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 1382-4 | 8.3 | 36 |
| 28 | Antiviral enantiomeric preference for 5Pnoraristeromycin. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 551-8.3 | 8.3 | 81 |
| 27 | 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 |
| 26 | Synthesis and antiviral activity of three pyrazole analogues of distamycin A. <i>Acta Chemica Scandinavica</i> , 1994 , 48, 498-505 | | 23 |

| | | | |
|----|---|------|-----|
| 25 | Inhibitory activity of S-adenosylhomocysteine hydrolase inhibitors against human cytomegalovirus replication. <i>Antiviral Research</i> , 1993 , 21, 197-216 | 10.8 | 39 |
| 24 | Molecular approaches for the treatment of hemorrhagic fever virus infections. <i>Antiviral Research</i> , 1993 , 22, 45-75 | 10.8 | 90 |
| 23 | Synthesis and antiviral activity of 5Pdeoxyypyrazofurin. <i>Journal of Medicinal Chemistry</i> , 1993 , 36, 3727-30 | 8.3 | 23 |
| 22 | Meningoradiculoneuritis due to acyclovir-resistant varicella-zoster virus in a patient with AIDS. <i>Journal of Infectious Diseases</i> , 1993 , 168, 1330-1 | 7 | 21 |
| 21 | S-Adenosyl-L-homocysteine Hydrolase Inhibitors as Anti-Viral Agents: 5?-Deoxyaristeromycin. <i>Nucleosides & Nucleotides</i> , 1993 , 12, 185-198 | | 26 |
| 20 | 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 |
| 19 | (E)-7-deazaaristeromycin lacking the hydroxymethyl substituent. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1993 , 3, 663-666 | 2.9 | 8 |
| 18 | Cyclopentene carbocyclic nucleosides related to the antitumor nucleoside clitocine and their conversion to 8-Aza-neplanocin analogues. Synthesis and antiviral activity. <i>Journal of Heterocyclic Chemistry</i> , 1993 , 30, 1393-1398 | 1.9 | 8 |
| 17 | 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 |
| 16 | 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 |
| 15 | Flow cytometric method for the detection of gpl antigens of varicella zoster virus and evaluation of anti-VZV agents. <i>Journal of Virological Methods</i> , 1992 , 38, 243-54 | 2.6 | 21 |
| 14 | 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 |
| 13 | Partially purified leaf extracts of <i>Melia azedarach</i> L. inhibit tacaribe virus growth in neonatal mice. <i>Phytotherapy Research</i> , 1992 , 6, 15-19 | 6.7 | 21 |
| 12 | 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 |
| 11 | 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 |
| 10 | 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 |
| 9 | 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 |
| 8 | 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 |

| | | | |
|---|---|------|----|
| 7 | 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 |
| 6 | Meliacine, an antiviral compound from <i>Melia azedarach</i> L., inhibits interferon production. <i>Journal of Interferon Research</i> , 1990 , 10, 469-75 | | 19 |
| 5 | 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 |
| 4 | Inhibitory effect of selected antiviral compounds on arenavirus replication in vitro. <i>Antiviral Research</i> , 1990 , 14, 287-99 | 10.8 | 88 |
| 3 | The antioxidizing effect of sterically hindered amines in thermal oxidation of low density polyethylene. <i>European Polymer Journal</i> , 1988 , 24, 289-294 | 5.2 | 10 |
| 2 | Induction of a refractory state to viral infection in mammalian cells by a plant inhibitor isolated from leaves of <i>Melia azedarach</i> L. <i>Antiviral Research</i> , 1988 , 9, 221-31 | 10.8 | 13 |
| 1 | An antiviral factor from <i>Melia azedarach</i> L. prevents Tacaribe virus encephalitis in mice. <i>Experientia</i> , 1986 , 42, 843-5 | | 23 |