

Dominique Schols

List of Publications by Year in descending order

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Version: 2024-02-01

431
papers

22,852
citations

11235

73
h-index

14386

132
g-index

467
all docs

467
docs citations

467
times ranked

20787
citing authors

#	ARTICLE	IF	CITATIONS
1	Itaconic acid hybrids as potential anticancer agents. <i>Molecular Diversity</i> , 2022, 26, 1-14.	2.1	9
2	Discovery of (±)-3-(1H-pyrazol-1-yl)-6,7-dihydro-5H-[1,2,4]triazolo[3,4-b][1,3,4] thiadiazine derivatives with promising in vitro anticoronavirus and antitumoral activity. <i>Molecular Diversity</i> , 2022, 26, 1357-1371.	2.1	11
3	Lung Microenvironments and Disease Progression in Fibrotic Hypersensitivity Pneumonitis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2022, 205, 60-74.	2.5	17
4	Atypical response to bacterial coinfection and persistent neutrophilic bronchoalveolar inflammation distinguish critical COVID-19 from influenza. <i>JCI Insight</i> , 2022, 7, .	2.3	38
5	Anthranilamides with quinoline and $\hat{1}^2$ -carboline scaffolds: design, synthesis, and biological activity. <i>Molecular Diversity</i> , 2022, 26, 2595-2612.	2.1	3
6	Synthesis, in vitro cytotoxicity, molecular docking and ADME study of some indolin-2-one linked 1,2,3-triazole derivatives. <i>Computational Biology and Chemistry</i> , 2022, 97, 107641.	1.1	4
7	Targeting chemokine receptors from the inside-out: discovery and development of small-molecule intracellular antagonists. <i>Chemical Communications</i> , 2022, 58, 4132-4148.	2.2	7
8	Innate Lymphoid Cells Are Required to Induce Airway Hyperreactivity in a Murine Neutrophilic Asthma Model. <i>Frontiers in Immunology</i> , 2022, 13, 849155.	2.2	7
9	Identification of novel chemotypes as CXCR2 antagonists via a scaffold hopping approach from a thiazolo[4,5-d]pyrimidine. <i>European Journal of Medicinal Chemistry</i> , 2022, 235, 114268.	2.6	7
10	In silico design, synthesis and anti-HIV activity of quinoline derivatives as non-nucleoside reverse transcriptase inhibitors (NNRTIs). <i>Computational Biology and Chemistry</i> , 2022, 98, 107675.	1.1	6
11	A Set of Experimentally Validated Decoys for the Human CC Chemokine Receptor 7 (CCR7) Obtained by Virtual Screening. <i>Frontiers in Pharmacology</i> , 2022, 13, 855653.	1.6	2
12	Effect of Particle Carriers for Intraperitoneal Drug Delivery on the Course of Ovarian Cancer and Its Immune Microenvironment in a Mouse Model. <i>Pharmaceutics</i> , 2022, 14, 687.	2.0	4
13	Discovery of 2-Phenylquinolines with Broad-Spectrum Anti-coronavirus Activity. <i>ACS Medicinal Chemistry Letters</i> , 2022, 13, 855-864.	1.3	10
14	SARS-CoV-2 Virion Infectivity and Cytokine Production in Primary Human Airway Epithelial Cells. <i>Viruses</i> , 2022, 14, 951.	1.5	6
15	Stimulation of the atypical chemokine receptor 3 (ACKR3) by a small-molecule agonist attenuates fibrosis in a preclinical liver but not lung injury model. <i>Cellular and Molecular Life Sciences</i> , 2022, 79, 293.	2.4	1
16	Organotropic dendrons with high potency as HIV-1, HIV-2 and EV-A71 cell entry inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2022, 237, 114414.	2.6	1
17	The discovery of Zika virus NS2B-NS3 inhibitors with antiviral activity via an integrated virtual screening approach. <i>European Journal of Pharmaceutical Sciences</i> , 2022, 175, 106220.	1.9	7
18	New 2-alkylthio-1-benzylimidazole-5-carboxylic acid derivatives targeting gp41: design, synthesis and in vitro anti-HIV activity evaluation. <i>Current HIV Research</i> , 2022, 20, .	0.2	0

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19	Potent neutralizing anti-SARS-CoV-2 human antibodies cure infection with SARS-CoV-2 variants in hamster model. <i>IScience</i> , 2022, 25, 104705.	1.9	8
20	β-Non-symmetrically Dimasked Tri- <i>o</i> -PPP Prodrugs as Potential Antiviral Agents against HIV. <i>ChemMedChem</i> , 2021, 16, 499-512.	1.6	15
21	Synthesis, Molecular Docking and Preliminary Antileukemic Activity of 4-Methoxybenzyl Derivatives Bearing Imidazo[2,1- <i>b</i>][1,3,4]thiadiazole. <i>Chemistry and Biodiversity</i> , 2021, 18, e2000800.	1.0	3
22	Palladium-catalyzed cross-coupling reactions on a bromo-naphthalene scaffold in the search for novel human CC chemokine receptor 8 (CCR8) antagonists. <i>Bioorganic Chemistry</i> , 2021, 107, 104560.	2.0	2
23	<i>In Vitro</i> Characterization of the Carbohydrate-Binding Agents HHA, GNA, and UDA as Inhibitors of Influenza A and B Virus Replication. <i>Antimicrobial Agents and Chemotherapy</i> , 2021, 65, .	1.4	8
24	Synthesis, in silico ADME, molecular docking and in vitro cytotoxicity evaluation of stilbene linked 1,2,3-triazoles. <i>Heliyon</i> , 2021, 7, e05893.	1.4	3
25	Targeted disruption of π-π stacking in Malaysian banana lectin reduces mitogenicity while preserving antiviral activity. <i>Scientific Reports</i> , 2021, 11, 656.	1.6	16
26	FO-SPR biosensor calibrated with recombinant extracellular vesicles enables specific and sensitive detection directly in complex matrices. <i>Journal of Extracellular Vesicles</i> , 2021, 10, e12059.	5.5	10
27	A novel experimental porcine model to assess the impact of differential pulmonary blood flow on ischemia-reperfusion injury after unilateral lung transplantation. <i>Intensive Care Medicine Experimental</i> , 2021, 9, 4.	0.9	5
28	Discovery of 3-phenyl- and 3-N-piperidinyl-isothiazolo[4,3- <i>b</i>]pyridines as highly potent inhibitors of cyclin G-associated kinase. <i>European Journal of Medicinal Chemistry</i> , 2021, 213, 113158.	2.6	10
29	Small Molecule Cyclotriazadisulfonamide Abrogates the Upregulation of the Human Receptors CD4 and 4-1BB and Suppresses In Vitro Activation and Proliferation of T Lymphocytes. <i>Frontiers in Immunology</i> , 2021, 12, 650731.	2.2	6
30	A patent review of adaptor associated kinase 1 (AAK1) inhibitors (2013-present). <i>Expert Opinion on Therapeutic Patents</i> , 2021, 31, 911-936.	2.4	15
31	Active Components from <i>Cassia abbreviata</i> Prevent HIV-1 Entry by Distinct Mechanisms of Action. <i>International Journal of Molecular Sciences</i> , 2021, 22, 5052.	1.8	6
32	Biological characterization of ligands targeting the human CC chemokine receptor 8 (CCR8) reveals the biased signaling properties of small molecule agonists. <i>Biochemical Pharmacology</i> , 2021, 188, 114565.	2.0	7
33	Synthesis, molecular docking, and preliminary cytotoxicity study of some novel 2-(naphthalen-1-yl)-methylimidazo[2,1- <i>b</i>][1,3,4]thiadiazoles. <i>Journal of Molecular Structure</i> , 2021, 1234, 130174.	1.8	2
34	Multivalent Tryptophan- and Tyrosine-Containing [60]Fullerene Hexa-Adducts as Dual HIV and Enterovirus A71 Entry Inhibitors. <i>Chemistry - A European Journal</i> , 2021, 27, 10700-10710.	1.7	9
35	A Novel Series of Indole Alkaloid Derivatives Inhibit Dengue and Zika Virus Infection by Interference with the Viral Replication Complex. <i>Antimicrobial Agents and Chemotherapy</i> , 2021, 65, e0234920.	1.4	13
36	Novel N-Substituted 3-Aryl-4-(diethoxyphosphoryl)azetidines as Antibiotic Enhancers and Antiviral Agents in Search for a Successful Treatment of Complex Infections. <i>International Journal of Molecular Sciences</i> , 2021, 22, 8032.	1.8	5

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37	Double Arylation of the Indole Side Chain of Tri- and Tetrapodal Tryptophan Derivatives Renders Highly Potent HIV-1 and EV-A71 Entry Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 10027-10046.	2.9	7
38	Water-Promoted Reaction of C ₆₀ /Ar ₅ Cl Compounds with Thiophenes Delivers a Family of Multifunctional Fullerene Derivatives with Selective Antiviral Properties. <i>Organic Letters</i> , 2021, 23, 7226-7230.	2.4	7
39	Broad spectrum anti-coronavirus activity of a series of anti-malaria quinoline analogues. <i>Antiviral Research</i> , 2021, 193, 105127.	1.9	27
40	Labyrinthopeptin A1 inhibits dengue and Zika virus infection by interfering with the viral phospholipid membrane. <i>Virology</i> , 2021, 562, 74-86.	1.1	12
41	Thiophene-2-carboxamide derivatives of anthraquinone: A new potent antitumor chemotype. <i>European Journal of Medicinal Chemistry</i> , 2021, 221, 113521.	2.6	12
42	Exploring the dNTP-binding site of HIV-1 reverse transcriptase for inhibitor design. <i>European Journal of Medicinal Chemistry</i> , 2021, 225, 113785.	2.6	3
43	Synthesis, Molecular Docking and Molecular Dynamics Simulation of 2-Thioxothiazolidin-4-One Derivatives against Gp41. <i>Current HIV Research</i> , 2021, 19, 47-60.	0.2	2
44	A Proteomic Study on the Membrane Protein Fraction of T Cells Confirms High Substrate Selectivity for the ER Translocation Inhibitor Cyclotriazadisulfonamide. <i>Molecular and Cellular Proteomics</i> , 2021, 20, 100144.	2.5	7
45	A single-dose live-attenuated YF17D-vectored SARS-CoV-2 vaccine candidate. <i>Nature</i> , 2021, 590, 320-325.	13.7	148
46	D-Peptide-Based Probe for CXCR4-Targeted Molecular Imaging and Radionuclide Therapy. <i>Pharmaceutics</i> , 2021, 13, 1619.	2.0	5
47	Synthesis and Anti-HIV Activity of a Novel Series of Isoquinoline-Based CXCR4 Antagonists. <i>Molecules</i> , 2021, 26, 6297.	1.7	2
48	Peripherally-driven myeloid NFκB and IFN/ISG responses predict malignancy risk, survival, and immunotherapy regime in ovarian cancer. , 2021, 9, e003609.		24
49	Skeleton binding protein-1-mediated parasite sequestration inhibits spontaneous resolution of malaria-associated acute respiratory distress syndrome. <i>PLoS Pathogens</i> , 2021, 17, e1010114.	2.1	7
50	Deciphering the Role of Extracellular Vesicles Derived from ZIKV-Infected hcMEC/D3 Cells on the Blood-Brain Barrier System. <i>Viruses</i> , 2021, 13, 2363.	1.5	8
51	Neo-Adjuvant Chemotherapy Reduces, and Surgery Increases Immunosuppression in First-Line Treatment for Ovarian Cancer. <i>Cancers</i> , 2021, 13, 5899.	1.7	9
52	Sliding of HIV-1 reverse transcriptase over DNA creates a transient P pocket “targeting P-pocket by fragment screening. <i>Nature Communications</i> , 2021, 12, 7127.	5.8	6
53	⁶⁴ Cu PET Imaging of the CXCR4 Chemokine Receptor Using a Cross-Bridged Cyclam Bis-Tetraazamacrocyclic Antagonist. <i>Journal of Nuclear Medicine</i> , 2020, 61, 123-128.	2.8	18
54	Design, synthesis and antiviral evaluation of novel acyclic phosphonate nucleotide analogs with triazolo[4,5- <i>b</i>]pyridine, imidazo[4,5- <i>b</i>]pyridine and imidazo[4,5- <i>b</i>]pyridin-2(3- <i>H</i>)-one systems. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2020, 39, 542-591.	0.4	8

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55	Synthesis, characterization and anti-HIV activity of polycarboxylic [60]fullerene derivatives obtained in the reaction of C ₆₀ Cl ₆ with a hydroquinone ether. <i>Tetrahedron Letters</i> , 2020, 61, 151598.	0.7	11
56	Direct arylation of C ₆₀ Cl ₆ and C ₇₀ Cl ₈ with carboxylic acids: a synthetic avenue to water-soluble fullerene derivatives with promising antiviral activity. <i>Chemical Communications</i> , 2020, 56, 1179-1182.	2.2	23
57	Labyrinthopeptins Exert Broad-Spectrum Antiviral Activity through Lipid-Binding-Mediated Virolysis. <i>Journal of Virology</i> , 2020, 94, .	1.5	30
58	Scaffold Simplification Strategy Leads to a Novel Generation of Dual Human Immunodeficiency Virus and Enterovirus-A71 Entry Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 349-368.	2.9	20
59	Preprotein signature for full susceptibility to the co-translational translocation inhibitor cyclotriazadisulfonamide. <i>Traffic</i> , 2020, 21, 250-264.	1.3	12
60	Assessment of protein biomarkers for preoperative differential diagnosis between benign and malignant ovarian tumors. <i>Gynecologic Oncology</i> , 2020, 159, 811-819.	0.6	8
61	Synthesis of new riboflavin modified ODNs: Effect of riboflavin moiety on the G-quadruplex arrangement and stability. <i>Bioorganic Chemistry</i> , 2020, 104, 104213.	2.0	0
62	Discovery of HIV entry inhibitors via a hybrid CXCR4 and CCR5 receptor pharmacophore-based virtual screening approach. <i>European Journal of Pharmaceutical Sciences</i> , 2020, 155, 105537.	1.9	22
63	Membrane Permeable, Bioreversibly Modified Prodrugs of Nucleoside Diphosphate- $\hat{3}$ -Phosphonates. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 11990-12007.	2.9	15
64	Thiophene-based water-soluble fullerene derivatives as highly potent antiherpetic pharmaceuticals. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 8702-8708.	1.5	4
65	Alkylated benzimidazoles: Design, synthesis, docking, DFT analysis, ADMET property, molecular dynamics and activity against HIV and YFV. <i>Computational Biology and Chemistry</i> , 2020, 89, 107400.	1.1	22
66	STAT2 signaling restricts viral dissemination but drives severe pneumonia in SARS-CoV-2 infected hamsters. <i>Nature Communications</i> , 2020, 11, 5838.	5.8	225
67	$\hat{3}$ -Ketobenzyl-Modified Nucleoside Triphosphate Prodrugs as Potential Antivirals. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 13745-13761.	2.9	10
68	Water-soluble fullerene-based nanostructures with promising antiviral and myogenic activity. <i>Chemical Communications</i> , 2020, 56, 10203-10206.	2.2	13
69	Development of a Novel SPR Assay to Study CXCR4-Ligand Interactions. <i>Biosensors</i> , 2020, 10, 150.	2.3	8
70	Anti-HIV-Active Nucleoside Triphosphate Prodrugs. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 6003-6027.	2.9	28
71	Prodrugs of $\hat{3}$ -Alkyl-Modified Nucleoside Triphosphates: Improved Inhibition of HIV Reverse Transcriptase. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 22063-22071.	7.2	19
72	Prodrugs of $\hat{3}$ -Alkyl-Modified Nucleoside Triphosphates: Improved Inhibition of HIV Reverse Transcriptase. <i>Angewandte Chemie</i> , 2020, 132, 22247-22255.	1.6	0

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73	Lipophilic Triphosphate Prodrugs of Various Nucleoside Analogues. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 6991-7007.	2.9	14
74	A chimeric yellow fever-Zika virus vaccine candidate fully protects against yellow fever virus infection in mice. <i>Emerging Microbes and Infections</i> , 2020, 9, 520-533.	3.0	21
75	Synthesis of 2-[(1 α -phthalimidoalkyl)sulfanyl]-pyrimidin-4(3H)-ones, their cytotoxicity and in vitro activity against HIV-1/2. <i>Chemistry of Heterocyclic Compounds</i> , 2020, 56, 67-72.	0.6	5
76	Design, synthesis and biological evaluation of 2-alkoxycarbonyl-3-anilinoindoles as a new class of potent inhibitors of tubulin polymerization. <i>Bioorganic Chemistry</i> , 2020, 97, 103665.	2.0	16
77	Early protein expression profile in bronchoalveolar lavage fluid and clinical outcomes in primary graft dysfunction after lung transplantation. <i>European Journal of Cardio-thoracic Surgery</i> , 2020, 58, 379-388.	0.6	4
78	Tryptophan Trimers and Tetramers Inhibit Dengue and Zika Virus Replication by Interfering with Viral Attachment Processes. <i>Antimicrobial Agents and Chemotherapy</i> , 2020, 64, .	1.4	8
79	Amides of pyrrole- and thiophene-fused anthraquinone derivatives: A role of the heterocyclic core in antitumor properties. <i>European Journal of Medicinal Chemistry</i> , 2020, 199, 112294.	2.6	22
80	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.	0.7	6
81	Synthesis and Evaluations of 1,4-Triazolyl Combretacoumarins and Desmethoxy Analogs. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 5610-5623.	1.2	7
82	A unique class of lignin derivatives displays broad anti-HIV activity by interacting with the viral envelope. <i>Virus Research</i> , 2019, 274, 197760.	1.1	10
83	Novel Isoxazolidine and β -Lactam Analogues of Homonucleosides. <i>Molecules</i> , 2019, 24, 4014.	1.7	14
84	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.	2.6	25
85	Acetate as a model for aspartate-based CXCR4 chemokine receptor binding of cobalt and nickel complexes of cross-bridged tetraazamacrocycles. <i>Dalton Transactions</i> , 2019, 48, 2785-2801.	1.6	11
86	Modifications in the branched arms of a class of dual inhibitors of HIV and EV71 replication expand their antiviral spectrum. <i>Antiviral Research</i> , 2019, 168, 210-214.	1.9	9
87	Diversion of the Arbuzov reaction: alkylation of C-Cl instead of phosphonic ester formation on the fullerene cage. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 7155-7160.	1.5	12
88	Advantages and shortcomings of cell-based electrical impedance measurements as a GPCR drug discovery tool. <i>Biosensors and Bioelectronics</i> , 2019, 137, 33-44.	5.3	20
89	Acyclic nucleoside phosphonates containing the amide bond: hydroxy derivatives. <i>Monatshefte für Chemie</i> , 2019, 150, 733-745.	0.9	2
90	Prone Positioning During Ex Vivo Lung Perfusion Influences Regional Edema Accumulation. <i>Journal of Surgical Research</i> , 2019, 239, 300-308.	0.8	24

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91	Highly stable hexitol based XNA aptamers targeting the vascular endothelial growth factor. <i>Nucleic Acids Research</i> , 2019, 47, 4927-4939.	6.5	73
92	Synthesis and Anti-HIV Activity of Guanine Modified Fluorinated Acyclic Nucleoside Phosphonate Derivatives. <i>Chemistry and Biodiversity</i> , 2019, 16, e1800532.	1.0	3
93	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.	0.7	7
94	CXCR7/ACKR3-targeting ligands interfere with X7 HIV-1 and HIV-2 entry and replication in human host cells. <i>Heliyon</i> , 2018, 4, e00557.	1.4	10
95	Synthesis of Enantiomerically Pure 1 β ,2 β -cis-dideoxy, -dideoxydi β -dehydro, -ribo and -deoxy Carbocyclic Nucleoside Analogues. <i>Synthesis</i> , 2018, 50, 2266-2280.	1.2	4
96	New antitumor anthra[2,3-b]furan-3-carboxamides: Synthesis and structure-activity relationship. <i>European Journal of Medicinal Chemistry</i> , 2018, 148, 128-139.	2.6	26
97	Different contributions of chemokine N β -terminal features attest to a different ligand binding mode and a bias towards activation of ACKR3/CXCR7 compared with CXCR4 and CXCR3. <i>British Journal of Pharmacology</i> , 2018, 175, 1419-1438.	2.7	52
98	A Flow Cytometry-based Assay to Identify Compounds That Disrupt Binding of Fluorescently-labeled CXC Chemokine Ligand 12 to CXC Chemokine Receptor 4. <i>Journal of Visualized Experiments</i> , 2018, , .	0.2	11
99	Aminomethylation of heliomycin: Preparation and anticancer characterization of the first series of semi-synthetic derivatives. <i>European Journal of Medicinal Chemistry</i> , 2018, 143, 1553-1562.	2.6	13
100	A yellow fever β -Zika chimeric virus vaccine candidate protects against Zika infection and congenital malformations in mice. <i>Npj Vaccines</i> , 2018, 3, 56.	2.9	41
101	Peroxynitrite Exposure of CXCL12 Impairs Monocyte, Lymphocyte and Endothelial Cell Chemotaxis, Lymphocyte Extravasation in vivo and Anti-HIV-1 Activity. <i>Frontiers in Immunology</i> , 2018, 9, 1933.	2.2	5
102	Tri-armed ligands of G-quadruplex on heteroarene-fused anthraquinone scaffolds: Design, synthesis and pre-screening of biological properties. <i>European Journal of Medicinal Chemistry</i> , 2018, 159, 59-73.	2.6	20
103	CXCR4-targeting nanobodies differentially inhibit CXCR4 function and HIV entry. <i>Biochemical Pharmacology</i> , 2018, 158, 402-412.	2.0	34
104	Nanobody-Fc constructs targeting chemokine receptor CXCR4 potently inhibit signaling and CXCR4-mediated HIV-entry and induce antibody effector functions. <i>Biochemical Pharmacology</i> , 2018, 158, 413-424.	2.0	44
105	Isoxazolidine Conjugates of N3-Substituted 6-Bromoquinazolinones β -Synthesis, Anti-Varizella-Zoster Virus, and Anti-Cytomegalovirus Activity. <i>Molecules</i> , 2018, 23, 1889.	1.7	10
106	Synthesis and Antiviral Activity of Water-Soluble Polycarboxylic Derivatives of [60]Fullerene Loaded with 3,4-Dichlorophenyl Units. <i>Chemistry and Biodiversity</i> , 2018, 15, e1800293.	1.0	7
107	A Kinetic Fluorescence-based Ca ²⁺ Mobilization Assay to Identify G Protein-coupled Receptor Agonists, Antagonists, and Allosteric Modulators. <i>Journal of Visualized Experiments</i> , 2018, , .	0.2	5
108	Expedient synthesis and biological evaluation of alkenyl acyclic nucleoside phosphonate prodrugs. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 3596-3609.	1.4	4

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109	Synthesis and Antiviral Evaluation of Tri(PPP)AbacavirTP, Tri(PPP)CarbovirTP, and Their 1,2,4-Disubstituted Analogues. <i>ChemMedChem</i> , 2018, 13, 1771-1778.	1.6	8
110	Asymmetric Primaquine and Halogenaniline Fumardiamides as Novel Biologically Active Michael Acceptors. <i>Molecules</i> , 2018, 23, 1724.	1.7	8
111	Synthesis of a 3-C-ethynyl- β -d-ribofuranose purine nucleoside library: Discovery of C7-deazapurine analogs as potent antiproliferative nucleosides. <i>European Journal of Medicinal Chemistry</i> , 2018, 157, 248-267.	2.6	15
112	Engineering <i>Lactobacillus rhamnosus</i> GG and GR-1 to express HIV-inhibiting griffithsin. <i>International Journal of Antimicrobial Agents</i> , 2018, 52, 599-607.	1.1	18
113	Iterative Chemical Engineering of Vancomycin Leads to Novel Vancomycin Analogs With a High in Vitro Therapeutic Index. <i>Frontiers in Microbiology</i> , 2018, 9, 1175.	1.5	9
114	Chloro-1,4-dimethyl-9H-carbazole Derivatives Displaying Anti-HIV Activity. <i>Molecules</i> , 2018, 23, 286.	1.7	15
115	Polyfunctionalized Pyrrole Derivatives: Easy Three-component Microwave-assisted Synthesis, Cytostatic and Antiviral Evaluation. <i>Current Microwave Chemistry</i> , 2018, 5, 23-31.	0.2	8
116	Design, synthesis, <i>in vitro</i> antiproliferative activity and apoptosis-inducing studies of 1-(3,4,5-trimethoxyphenyl)-3-(2-alkoxycarbonylindolyl)-2-propen-1-one derivatives obtained by a molecular hybridisation approach. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018, 33, 1225-1238.	2.5	16
117	Facile Microwave-assisted Synthesis of Various C5-modified Pyrimidine Pyranonucleosides as Potential Cytotoxic Antitumor Agents. <i>Current Microwave Chemistry</i> , 2018, 4, .	0.2	1
118	A Proteomic Survey Indicates Sortilin as a Secondary Substrate of the ER Translocation Inhibitor Cyclotriazadisulfonamide (CADA). <i>Molecular and Cellular Proteomics</i> , 2017, 16, 157-167.	2.5	17
119	Structure-activity relationship studies on a Trp dendrimer with dual activities against HIV and enterovirus A71. Modifications on the amino acid. <i>Antiviral Research</i> , 2017, 139, 32-40.	1.9	17
120	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.	1.5	17
121	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.	2.1	8
122	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.	2.9	22
123	Synthesis of different types of alkoxy fullerene derivatives from chlorofullerene $C_{60}Cl_6$. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 773-777.	1.5	28
124	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.	2.6	27
125	Signaling properties of the human chemokine receptors CXCR4 and CXCR7 by cellular electric impedance measurements. <i>PLoS ONE</i> , 2017, 12, e0185354.	1.1	21
126	Comparison of cell-based assays for the identification and evaluation of competitive CXCR4 inhibitors. <i>PLoS ONE</i> , 2017, 12, e0176057.	1.1	33

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127	Immunoregulatory effects of multipotent adult progenitor cells in a porcine ex vivo lung perfusion model. <i>Stem Cell Research and Therapy</i> , 2017, 8, 159.	2.4	51
128	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.	0.7	9
129	A porcine ex vivo lung perfusion model with maximal argon exposure to attenuate ischemia-reperfusion injury. <i>Medical Gas Research</i> , 2017, 7, 28.	1.2	8
130	New Isoxazolidine-Conjugates of Quinazolinonesâ€™Synthesis, Antiviral and Cytostatic Activity. <i>Molecules</i> , 2016, 21, 959.	1.7	19
131	Design, Synthesis and Biological Evaluation of Novel Primaquine-Cinnamic Acid Conjugates of the Amide and Acylsemicarbazide Type. <i>Molecules</i> , 2016, 21, 1629.	1.7	22
132	Development and Identification of a Novel Anti-HIV-1 Peptide Derived by Modification of the N-Terminal Domain of HIV-1 Integrase. <i>Frontiers in Microbiology</i> , 2016, 7, 845.	1.5	13
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393	New Polyacetal Polysulphate Active against Human Immunodeficiency Virus and other Enveloped Viruses. <i>Antiviral Chemistry and Chemotherapy</i> , 1992, 3, 351-360.	0.3	19
394	Differential Activity of Polyanionic Compounds and Castanospermine against HIV Replication and HIV-Induced Syncytium Formation Depending on Virus Strain and Cell Type. <i>Antiviral Chemistry and Chemotherapy</i> , 1992, 3, 23-29.	0.3	25
395	2',5'-Bis-O-(tert-butylidimethylsilyl)-3'-spiro-5''-(4''-amino-1'',2''-oxathiole-2'',2'-dioxide)pyrimidine (TSAO) nucleoside analogues: highlyselective inhibitors of human immunodeficiency virus type 1 that are targeted at the viral reverse transcriptase.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1992, 89, 4392-4396.	3.3	164
396	Sulfonic acid polymers as a new class of human immunodeficiency virus inhibitors. <i>Antiviral Research</i> , 1992, 18, 139-150.	1.9	68

#	ARTICLE	IF	CITATIONS
397	The mannose-specific plant lectins from <i>Cymbidium hybrid</i> and <i>Epipactis helleborine</i> and the (N-acetylglucosamine)n-specific plant lectin from <i>Urtica dioica</i> are potent and selective inhibitors of human immunodeficiency virus and cytomegalovirus replication in vitro. <i>Antiviral Research</i> , 1992, 18, 191-207.	1.9	230
398	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, 243-254.	1.0	23
399	Structure investigation and anti-HIV activities of high-molecular weight ATA polymers. <i>Journal of Organic Chemistry</i> , 1992, 57, 7241-7248.	1.7	20
400	Anti-human immunodeficiency virus effects of cationic metalloporphyrin-ellipticine complexes. <i>Biochemical Pharmacology</i> , 1992, 44, 1675-1679.	2.0	45
401	Sulfated polymers inhibit the interaction of human cytomegalovirus with cell surface heparan sulfate. <i>Virology</i> , 1992, 189, 48-58.	1.1	173
402	Presence of class II histocompatibility DR proteins on the envelope of human immunodeficiency virus demonstrated by FACS analysis. <i>Virology</i> , 1992, 189, 374-376.	1.1	71
403	Activity of different antiviral drug combinations against human cytomegalovirus replication in vitro. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 1992, 11, 1144-1155.	1.3	31
404	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.	1.9	30
405	Preparation and anti-HIV activities of aurintricarboxylic acid fractions and analogs: direct correlation of antiviral potency with molecular weight. <i>Journal of Medicinal Chemistry</i> , 1991, 34, 329-337.	2.9	80
406	1- β -D-Ribofuranosyl-1,2,4-triazole-3-carboxamide (ribavirin) and 5-ethynyl-1- β -D-ribofuranosylimidazole-4-carboxamide (EICAR) markedly potentiate the inhibitory effect of 2',3'-dideoxyinosine on human immunodeficiency virus in peripheral blood lymphocytes. <i>Biochemical and Biophysical Research Communications</i> , 1991, 178, 563-569.	1.0	31
407	Activity of acyclic nucleoside phosphonate analogues against human immunodeficiency virus in monocyte/macrophages and peripheral blood lymphocytes. <i>Biochemical and Biophysical Research Communications</i> , 1991, 178, 329-335.	1.0	55
408	Sensitive, reproducible and convenient fluorometric assay for the in vitro evaluation of anticytomegalovirus agents. <i>Journal of Virological Methods</i> , 1991, 35, 27-38.	1.0	22
409	Synthesis and anti-HIV activities of low molecular weight aurintricarboxylic acid fragments and related compounds. <i>Journal of Medicinal Chemistry</i> , 1991, 34, 337-342.	2.9	61
410	Comparative activity of selected antiviral compounds against clinical isolates of human cytomegalovirus. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 1991, 10, 1026-1033.	1.3	68
411	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.	0.3	54
412	Sulphated Cyclodextrins are Potent anti-HIV Agents Acting Synergistically with 2',3'-dideoxynucleoside Analogues. <i>Antiviral Chemistry and Chemotherapy</i> , 1991, 2, 45-53.	0.3	41
413	Inhibitory Effects of Polycations on the Replication of Enveloped Viruses (HIV, HSV, CMV, RSV), Tj ETQq1 1 0.784314 rgBT /Overlock 10 243-248.	0.3	18
414	Potent and selective inhibition of HIV-1 replication in vitro by a novel series of TIBO derivatives. <i>Nature</i> , 1990, 343, 470-474.	13.7	794

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416	Dextran sulfate and other polyanionic anti-HIV compounds specifically interact with the viral gp120 glycoprotein expressed by T-cells persistently infected with HIV-1. <i>Virology</i> , 1990, 175, 556-561.	1.1	217
417	Flow cytometric method to monitor the destruction of CD4+ cells following their fusion with HIV-infected cells. <i>Cytometry</i> , 1990, 11, 736-743.	1.8	19
418	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.	0.3	85
419	Anti-HIV-1 activity of antiviral compounds, as quantitated by a focal immunoassay in CD4+ HeLa cells and a plaque assay in MT-4 cells. <i>Journal of Virological Methods</i> , 1990, 29, 197-208.	1.0	21
420	Syncytium Formation and Destruction of Bystander CD4+ Cells Cocultured with T Cells Persistently Infected with Human Immunodeficiency Virus as Demonstrated by Flow Cytometry. <i>Journal of General Virology</i> , 1989, 70, 2397-2408.	1.3	49
421	Detection of immediate early, early and late antigens of human cytomegalovirus by flow cytometry. <i>Journal of Virological Methods</i> , 1989, 26, 247-254.	1.0	32
422	Tetrazolium-based plaque assay for HIV-1 and HIV-2, and its use in the evaluation of antiviral compounds. <i>Journal of Virological Methods</i> , 1989, 26, 319-329.	1.0	36
423	Highly specific inhibition of human immunodeficiency virus type 1 by a novel 6-substituted acyclouridine derivative. <i>Biochemical and Biophysical Research Communications</i> , 1989, 165, 1375-1381.	1.0	294
424	Specific interaction of aurointricarboxylic acid with the human immunodeficiency virus/CD4 cell receptor.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1989, 86, 3322-3326.	3.3	118
425	A highly reliable, sensitive, flow cytometric/fluorometric assay for the evaluation of the anti-HIV activity of antiviral compounds in MT-4 cells. <i>Journal of Immunological Methods</i> , 1988, 114, 27-32.	0.6	28
426	Rapid and automated tetrazolium-based colorimetric assay for the detection of anti-HIV compounds. <i>Journal of Virological Methods</i> , 1988, 20, 309-321.	1.0	1,644
427	Fuchsin acid selectively inhibits human immunodeficiency virus (HIV) replication invitro. <i>Biochemical and Biophysical Research Communications</i> , 1988, 155, 1404-1411.	1.0	48
428	Pentosan polysulfate, a sulfated oligosaccharide, is a potent and selective anti-HIV agent in vitro. <i>Antiviral Research</i> , 1988, 9, 335-343.	1.9	195
429	Immunocytochemical demonstration of proopiomelanocortin- and other opioid-related substances and a CRF-like peptide in the gut of the american cockroach, <i>Periplaneta americana</i> L. <i>Histochemistry</i> , 1987, 86, 345-351.	1.9	38
430	Broad Antiviral Activity of Carbohydrate-Binding Agents Against Dengue Virus Infection. , 0, , .		0
431	In-Depth Characterization of Zika Virus Inhibitors Using Cell-Based Electrical Impedance. <i>Microbiology Spectrum</i> , 0, , .	1.2	4