## **Dominique Schols**

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Itaconic acid hybrids as potential anticancer agents. Molecular Diversity, 2022, 26, 1-14.	3.9	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. Molecular Diversity, 2022, 26, 1357-1371.	3.9	11
3	Lung Microenvironments and Disease Progression in Fibrotic Hypersensitivity Pneumonitis. American Journal of Respiratory and Critical Care Medicine, 2022, 205, 60-74.	5.6	17
4	Atypical response to bacterial coinfection and persistent neutrophilic bronchoalveolar inflammation distinguish critical COVID-19 from influenza. JCI Insight, 2022, 7, .	5.0	38
5	Anthranilamides with quinoline and β-carboline scaffolds: design, synthesis, and biological activity. Molecular Diversity, 2022, 26, 2595-2612.	3.9	3
6	Synthesis, in vitro cytotoxicity,Âmolecular docking and ADME study of some indolin-2-one linked 1,2,3-triazole derivatives. Computational Biology and Chemistry, 2022, 97, 107641.	2.3	4
7	Targeting chemokine receptors from the inside-out: discovery and development of small-molecule intracellular antagonists. Chemical Communications, 2022, 58, 4132-4148.	4.1	7
8	Innate Lymphoid Cells Are Required to Induce Airway Hyperreactivity in a Murine Neutrophilic Asthma Model. Frontiers in Immunology, 2022, 13, 849155.	4.8	7
9	Identification of novel chemotypes as CXCR2 antagonists via a scaffold hopping approach from a thiazolo[4,5-d]pyrimidine. European Journal of Medicinal Chemistry, 2022, 235, 114268.	5.5	7
10	In silico design, synthesis and anti-HIV activity of quinoline derivatives as non-nucleoside reverse transcriptase inhibitors (NNRTIs). Computational Biology and Chemistry, 2022, 98, 107675.	2.3	6
11	A Set of Experimentally Validated Decoys for the Human CC Chemokine Receptor 7 (CCR7) Obtained by Virtual Screening. Frontiers in Pharmacology, 2022, 13, 855653.	3.5	2
12	Effect of Particle Carriers for Intraperitoneal Drug Delivery on the Course of Ovarian Cancer and Its Immune Microenvironment in a Mouse Model. Pharmaceutics, 2022, 14, 687.	4.5	4
13	Discovery of 2-Phenylquinolines with Broad-Spectrum Anti-coronavirus Activity. ACS Medicinal Chemistry Letters, 2022, 13, 855-864.	2.8	10
14	SARS-CoV-2 Virion Infectivity and Cytokine Production in Primary Human Airway Epithelial Cells. Viruses, 2022, 14, 951.	3.3	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. Cellular and Molecular Life Sciences, 2022, 79, 293.	5.4	1
16	Organotropic dendrons with high potency as HIV-1, HIV-2 and EV-A71 cell entry inhibitors. European Journal of Medicinal Chemistry, 2022, 237, 114414.	5.5	1
17	The discovery of Zika virus NS2B-NS3 inhibitors with antiviral activity via an integrated virtual screening approach. European Journal of Pharmaceutical Sciences, 2022, 175, 106220.	4.0	7
18	New 2-alkylthio-1-benzylimidazole-5-carboxylic acid derivatives targeting gp41: design, synthesis and in vitro anti-HIV activity evaluation. Current HIV Research, 2022, 20, .	0.5	0

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19	Potent neutralizing anti-SARS-CoV-2 human antibodies cure infection with SARS-CoV-2 variants in hamster model. IScience, 2022, 25, 104705.	4.1	8
20	γâ€Nonâ€Symmetrically Dimasked Tri <i>PPP</i> ro Prodrugs as Potential Antiviral Agents against HIV. ChemMedChem, 2021, 16, 499-512.	3.2	15
21	Synthesis, Molecular Docking and Preliminary Antileukemic Activity of 4â€Methoxybenzyl Derivatives Bearing Imidazo[2,1―b ][1,3,4]thiadiazole. Chemistry and Biodiversity, 2021, 18, e2000800.	2.1	3
22	Palladium-catalyzed cross-coupling reactions on a bromo-naphthalene scaffold in the search for novel human CC chemokine receptor 8 (CCR8) antagonists. Bioorganic Chemistry, 2021, 107, 104560.	4.1	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. Antimicrobial Agents and Chemotherapy, 2021, 65, .	3.2	8
24	Synthesis, in silico ADME, molecular docking and in vitro cytotoxicity evaluation of stilbene linked 1,2,3-triazoles. Heliyon, 2021, 7, e05893.	3.2	3
25	Targeted disruption of pi–pi stacking in Malaysian banana lectin reduces mitogenicity while preserving antiviral activity. Scientific Reports, 2021, 11, 656.	3.3	16
26	FO‣PR biosensor calibrated with recombinant extracellular vesicles enables specific and sensitive detection directly in complex matrices. Journal of Extracellular Vesicles, 2021, 10, e12059.	12.2	10
27	A novel experimental porcine model to assess the impact of differential pulmonary blood flow on ischemia–reperfusion injury after unilateral lung transplantation. Intensive Care Medicine Experimental, 2021, 9, 4.	1.9	5
28	Discovery of 3-phenyl- and 3-N-piperidinyl-isothiazolo[4,3-b]pyridines as highly potent inhibitors of cyclin G-associated kinase. European Journal of Medicinal Chemistry, 2021, 213, 113158.	5.5	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. Frontiers in Immunology, 2021, 12, 650731.	4.8	6
30	A patent review of adaptor associated kinase 1 (AAK1) inhibitors (2013-present). Expert Opinion on Therapeutic Patents, 2021, 31, 911-936.	5.0	15
31	Active Components from Cassia abbreviata Prevent HIV-1 Entry by Distinct Mechanisms of Action. International Journal of Molecular Sciences, 2021, 22, 5052.	4.1	6
32	Biological characterization of ligands targeting the human CC chemokine receptor 8 (CCR8) reveals the biased signaling properties of small molecule agonists. Biochemical Pharmacology, 2021, 188, 114565.	4.4	7
33	Synthesis, molecular docking, and preliminary cytotoxicity study of some novel 2-(naphthalen-1-yl)-methylimidazo[2,1-b][1,3,4]thiadiazoles. Journal of Molecular Structure, 2021, 1234, 130174.	3.6	2
34	Multivalent Tryptophan―and Tyrosineâ€Containing [60]Fullerene Hexaâ€Adducts as Dual HIV and Enterovirus A71 Entry Inhibitors. Chemistry - A European Journal, 2021, 27, 10700-10710.	3.3	9
35	A Novel Series of Indole Alkaloid Derivatives Inhibit Dengue and Zika Virus Infection by Interference with the Viral Replication Complex. Antimicrobial Agents and Chemotherapy, 2021, 65, e0234920.	3.2	13
36	Novel N-Substituted 3-Aryl-4-(diethoxyphosphoryl)azetidin-2-ones as Antibiotic Enhancers and Antiviral Agents in Search for a Successful Treatment of Complex Infections. International Journal of Molecular Sciences, 2021, 22, 8032.	4.1	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. Journal of Medicinal Chemistry, 2021, 64, 10027-10046.	6.4	7
38	Water-Promoted Reaction of C <sub>60</sub> Ar <sub>5</sub> Cl Compounds with Thiophenes Delivers a Family of Multifunctional Fullerene Derivatives with Selective Antiviral Properties. Organic Letters, 2021, 23, 7226-7230.	4.6	7
39	Broad spectrum anti-coronavirus activity of a series of anti-malaria quinoline analogues. Antiviral Research, 2021, 193, 105127.	4.1	27
40	Labyrinthopeptin A1 inhibits dengue and Zika virus infection by interfering with the viral phospholipid membrane. Virology, 2021, 562, 74-86.	2.4	12
41	Thiophene-2-carboxamide derivatives of anthraquinone: A new potent antitumor chemotype. European Journal of Medicinal Chemistry, 2021, 221, 113521.	5.5	12
42	Exploring the dNTP -binding site of HIV-1 reverse transcriptase for inhibitor design. European Journal of Medicinal Chemistry, 2021, 225, 113785.	5.5	3
43	Synthesis, Molecular Docking and Molecular Dynamics Simulation of 2- Thioxothiazolidin-4-One Derivatives against Gp41. Current HIV Research, 2021, 19, 47-60.	0.5	2
44	A Proteomic Study on the Membrane Protein Fraction of T Cells Confirms High Substrate Selectivity for the ER Translocation Inhibitor Cyclotriazadisulfonamide. Molecular and Cellular Proteomics, 2021, 20, 100144.	3.8	7
45	A single-dose live-attenuated YF17D-vectored SARS-CoV-2 vaccine candidate. Nature, 2021, 590, 320-325.	27.8	148
46	D-Peptide-Based Probe for CXCR4-Targeted Molecular Imaging and Radionuclide Therapy. Pharmaceutics, 2021, 13, 1619.	4.5	5
47	Synthesis and Anti-HIV Activity of a Novel Series of Isoquinoline-Based CXCR4 Antagonists. Molecules, 2021, 26, 6297.	3.8	2
48	Peripherally-driven myeloid NFkB and IFN/ISC 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. PLoS Pathogens, 2021, 17, e1010114.	4.7	7
50	Deciphering the Role of Extracellular Vesicles Derived from ZIKV-Infected hcMEC/D3 Cells on the Blood–Brain Barrier System. Viruses, 2021, 13, 2363.	3.3	8
51	Neo-Adjuvant Chemotherapy Reduces, and Surgery Increases Immunosuppression in First-Line Treatment for Ovarian Cancer. Cancers, 2021, 13, 5899.	3.7	9
52	Sliding of HIV-1 reverse transcriptase over DNA creates a transient P pocket – targeting P-pocket by fragment screening. Nature Communications, 2021, 12, 7127.	12.8	6
53	<sup>64</sup> Cu PET Imaging of the CXCR4 Chemokine Receptor Using a Cross-Bridged Cyclam Bis-Tetraazamacrocyclic Antagonist. Journal of Nuclear Medicine, 2020, 61, 123-128.	5.0	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. Nucleosides, Nucleotides and Nucleic Acids, 2020, 39, 542-591.	1.1	8

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55	Synthesis, characterization and anti-HIV activity of polycarboxylic [60]fullerene derivatives obtained in the reaction of C60Cl6 with a hydroquinone ether. Tetrahedron Letters, 2020, 61, 151598.	1.4	11
56	Direct arylation of C <sub>60</sub> Cl <sub>6</sub> and C <sub>70</sub> Cl <sub>8</sub> with carboxylic acids: a synthetic avenue to water-soluble fullerene derivatives with promising antiviral activity. Chemical Communications, 2020, 56, 1179-1182.	4.1	23
57	Labyrinthopeptins Exert Broad-Spectrum Antiviral Activity through Lipid-Binding-Mediated Virolysis. Journal of Virology, 2020, 94, .	3.4	30
58	Scaffold Simplification Strategy Leads to a Novel Generation of Dual Human Immunodeficiency Virus and Enterovirus-A71 Entry Inhibitors. Journal of Medicinal Chemistry, 2020, 63, 349-368.	6.4	20
59	Preprotein signature for full susceptibility to the coâ€ŧranslational translocation inhibitor cyclotriazadisulfonamide. Traffic, 2020, 21, 250-264.	2.7	12
60	Assessment of protein biomarkers for preoperative differential diagnosis between benign and malignant ovarian tumors. Gynecologic Oncology, 2020, 159, 811-819.	1.4	8
61	Synthesis of new riboflavin modified ODNs: Effect of riboflavin moiety on the G-quadruplex arrangement and stability. Bioorganic Chemistry, 2020, 104, 104213.	4.1	0
62	Discovery of HIV entry inhibitors via a hybrid CXCR4 and CCR5 receptor pharmacophoreâ€based virtual screening approach. European Journal of Pharmaceutical Sciences, 2020, 155, 105537.	4.0	22
63	Membrane Permeable, Bioreversibly Modified Prodrugs of Nucleoside Diphosphate-Î <sup>3</sup> -Phosphonates. Journal of Medicinal Chemistry, 2020, 63, 11990-12007.	6.4	15
64	Thiophene-based water-soluble fullerene derivatives as highly potent antiherpetic pharmaceuticals. Organic and Biomolecular Chemistry, 2020, 18, 8702-8708.	2.8	4
65	Alkylated benzimidazoles: Design, synthesis, docking, DFT analysis, ADMET property, molecular dynamics and activity against HIV and YFV. Computational Biology and Chemistry, 2020, 89, 107400.	2.3	22
66	STAT2 signaling restricts viral dissemination but drives severe pneumonia in SARS-CoV-2 infected hamsters. Nature Communications, 2020, 11, 5838.	12.8	225
67	Î <sup>3</sup> -Ketobenzyl-Modified Nucleoside Triphosphate Prodrugs as Potential Antivirals. Journal of Medicinal Chemistry, 2020, 63, 13745-13761.	6.4	10
68	Water-soluble fullerene-based nanostructures with promising antiviral and myogenic activity. Chemical Communications, 2020, 56, 10203-10206.	4.1	13
69	Development of a Novel SPR Assay to Study CXCR4–Ligand Interactions. Biosensors, 2020, 10, 150.	4.7	8
70	Anti-HIV-Active Nucleoside Triphosphate Prodrugs. Journal of Medicinal Chemistry, 2020, 63, 6003-6027.	6.4	28
71	Prodrugs of γâ€Alkylâ€Modified Nucleoside Triphosphates: Improved Inhibition of HIV Reverse Transcriptase. Angewandte Chemie - International Edition, 2020, 59, 22063-22071.	13.8	19
72	Prodrugs of γâ€Alkylâ€Modified Nucleoside Triphosphates: Improved Inhibition of HIV Reverse Transcriptase. Angewandte Chemie, 2020, 132, 22247-22255.	2.0	0

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73	Lipophilic Triphosphate Prodrugs of Various Nucleoside Analogues. Journal of Medicinal Chemistry, 2020, 63, 6991-7007.	6.4	14
74	A chimeric yellow fever-Zika virus vaccine candidate fully protects against yellow fever virus infection in mice. Emerging Microbes and Infections, 2020, 9, 520-533.	6.5	21
75	Synthesis of 2-[(ω-phthalimidoalkyl)sulfanyl]- pyrimidin-4(3H)-ones, their cytotoxicity and in vitro activity against HIV-1/2. Chemistry of Heterocyclic Compounds, 2020, 56, 67-72.	1.2	5
76	Design, synthesis and biological evaluation of 2-alkoxycarbonyl-3-anilinoindoles as a new class of potent inhibitors of tubulin polymerization. Bioorganic Chemistry, 2020, 97, 103665.	4.1	16
77	Early protein expression profile in bronchoalveolar lavage fluid and clinical outcomes in primary graft dysfunction after lung transplantation. European Journal of Cardio-thoracic Surgery, 2020, 58, 379-388.	1.4	4
78	Tryptophan Trimers and Tetramers Inhibit Dengue and Zika Virus Replication by Interfering with Viral Attachment Processes. Antimicrobial Agents and Chemotherapy, 2020, 64, .	3.2	8
79	Amides of pyrrole- and thiophene-fused anthraquinone derivatives: A role of the heterocyclic core in antitumor properties. European Journal of Medicinal Chemistry, 2020, 199, 112294.	5.5	22
80	Design, Synthesis, and Biological Evaluation of Novel C5-Modified Pyrimidine Ribofuranonucleosides as Potential Antitumor or/and Antiviral Agents. Medicinal Chemistry, 2020, 16, 368-384.	1.5	6
81	Synthesis and Evaluations of "1,4â€Triazolyl Combretacoumarins―and Desmethoxy Analogs. European Journal of Organic Chemistry, 2019, 2019, 5610-5623.	2.4	7
82	A unique class of lignin derivatives displays broad anti-HIV activity by interacting with the viral envelope. Virus Research, 2019, 274, 197760.	2.2	10
83	Novel Isoxazolidine and $\hat{I}^3$ -Lactam Analogues of Homonucleosides. Molecules, 2019, 24, 4014.	3.8	14
84	Antitumor and antiviral activities of 4-substituted 1,2,3-triazolyl-2,3-dibenzyl-L-ascorbic acid derivatives. European Journal of Medicinal Chemistry, 2019, 184, 111739.	5.5	25
85	Acetate as a model for aspartate-based CXCR4 chemokine receptor binding of cobalt and nickel complexes of cross-bridged tetraazamacrocycles. Dalton Transactions, 2019, 48, 2785-2801.	3.3	11
86	Modifications in the branched arms of a class of dual inhibitors of HIV and EV71 replication expand their antiviral spectrum. Antiviral Research, 2019, 168, 210-214.	4.1	9
87	Diversion of the Arbuzov reaction: alkylation of C–Cl instead of phosphonic ester formation on the fullerene cage. Organic and Biomolecular Chemistry, 2019, 17, 7155-7160.	2.8	12
88	Advantages and shortcomings of cell-based electrical impedance measurements as a GPCR drug discovery tool. Biosensors and Bioelectronics, 2019, 137, 33-44.	10.1	20
89	Acyclic nucleoside phosphonates containing the amide bond: hydroxy derivatives. Monatshefte Für Chemie, 2019, 150, 733-745.	1.8	2
90	Prone Positioning During ExÂVivo Lung Perfusion Influences Regional Edema Accumulation. Journal of Surgical Research, 2019, 239, 300-308.	1.6	24

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91	Highly stable hexitol based XNA aptamers targeting the vascular endothelial growth factor. Nucleic Acids Research, 2019, 47, 4927-4939.	14.5	73
92	Synthesis and Anti-HIV Activity of Guanine Modified Fluorinated Acyclic Nucleoside Phosphonate Derivatives. Chemistry and Biodiversity, 2019, 16, e1800532.	2.1	3
93	Synthesis, Anti-Varicella-Zoster Virus and Anti-Cytomegalovirus Activity of 4,5-Disubstituted 1,2,3-(1H)-Triazoles. Medicinal Chemistry, 2019, 15, 801-812.	1.5	7
94	CXCR7/ACKR3-targeting ligands interfere with X7 HIV-1 and HIV-2 entry and replication in human host cells. Heliyon, 2018, 4, e00557.	3.2	10
95	Synthesis of Enantiomerically Pure 1′,2′-cis-dideoxy, -dideoxydiÂdehydro, -ribo and -deoxy Carbocyclic Nucleoside Analogues. Synthesis, 2018, 50, 2266-2280.	2.3	4
96	New antitumor anthra[2,3-b]furan-3-carboxamides: Synthesis and structure-activity relationship. European Journal of Medicinal Chemistry, 2018, 148, 128-139.	5.5	26
97	Different contributions of chemokine Nâ€ŧerminal features attest to a different ligand binding mode and a bias towards activation of ACKR3/CXCR7 compared with CXCR4 and CXCR3. British Journal of Pharmacology, 2018, 175, 1419-1438.	5.4	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. Journal of Visualized Experiments, 2018, , .	0.3	11
99	Aminomethylation of heliomycin: Preparation and anticancer characterization of the first series of semi-synthetic derivatives. European Journal of Medicinal Chemistry, 2018, 143, 1553-1562.	5.5	13
100	A yellow fever–Zika chimeric virus vaccine candidate protects against Zika infection and congenital malformations in mice. Npj Vaccines, 2018, 3, 56.	6.0	41
101	Peroxynitrite Exposure of CXCL12 Impairs Monocyte, Lymphocyte and Endothelial Cell Chemotaxis, Lymphocyte Extravasation in vivo and Anti-HIV-1 Activity. Frontiers in Immunology, 2018, 9, 1933.	4.8	5
102	Tri-armed ligands of G-quadruplex on heteroarene-fused anthraquinone scaffolds: Design, synthesis and pre-screening of biological properties. European Journal of Medicinal Chemistry, 2018, 159, 59-73.	5.5	20
103	CXCR4-targeting nanobodies differentially inhibit CXCR4 function and HIV entry. Biochemical Pharmacology, 2018, 158, 402-412.	4.4	34
104	Nanobody-Fc constructs targeting chemokine receptor CXCR4 potently inhibit signaling and CXCR4-mediated HIV-entry and induce antibody effector functions. Biochemical Pharmacology, 2018, 158, 413-424.	4.4	44
105	Isoxazolidine Conjugates of N3-Substituted 6-Bromoquinazolinones—Synthesis, Anti-Varizella-Zoster Virus, and Anti-Cytomegalovirus Activity. Molecules, 2018, 23, 1889.	3.8	10
106	Synthesis and Antiviral Activity of Water-Soluble Polycarboxylic Derivatives of [60]Fullerene Loaded with 3,4-Dichlorophenyl Units. Chemistry and Biodiversity, 2018, 15, e1800293.	2.1	7
107	A Kinetic Fluorescence-based Ca <sup>2+</sup> Mobilization Assay to Identify G Protein-coupled Receptor Agonists, Antagonists, and Allosteric Modulators. Journal of Visualized Experiments, 2018, , .	0.3	5
108	Expedient synthesis and biological evaluation of alkenyl acyclic nucleoside phosphonate prodrugs. Bioorganic and Medicinal Chemistry, 2018, 26, 3596-3609.	3.0	4

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109	Synthesis and Antiviral Evaluation of Tri <i>PPP</i> roâ€AbacavirTP, Tri <i>PPP</i> roâ€CarbovirTP, and Their 1′,2′â€ <i>cis</i> â€Disubstituted Analogues. ChemMedChem, 2018, 13, 1771-1778.	3.2	8
110	Asymmetric Primaquine and Halogenaniline Fumardiamides as Novel Biologically Active Michael Acceptors. Molecules, 2018, 23, 1724.	3.8	8
111	Synthesis of a 3′-C-ethynyl-β-d-ribofuranose purine nucleoside library: Discovery of C7-deazapurine analogs as potent antiproliferative nucleosides. European Journal of Medicinal Chemistry, 2018, 157, 248-267.	5.5	15
112	Engineering Lactobacillus rhamnosus GG and GR-1 to express HIV-inhibiting griffithsin. International Journal of Antimicrobial Agents, 2018, 52, 599-607.	2.5	18
113	Iterative Chemical Engineering of Vancomycin Leads to Novel Vancomycin Analogs With a High in Vitro Therapeutic Index. Frontiers in Microbiology, 2018, 9, 1175.	3.5	9
114	Chloro-1,4-dimethyl-9H-carbazole Derivatives Displaying Anti-HIV Activity. Molecules, 2018, 23, 286.	3.8	15
115	Polyfunctionalized Pyrrole Derivatives: Easy Three-component Microwave-assisted Synthesis, Cytostatic and Antiviral Evaluation. Current Microwave Chemistry, 2018, 5, 23-31.	0.8	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. Journal of Enzyme Inhibition and Medicinal Chemistry, 2018, 33, 1225-1238	5.2	16
117	Facile Microwave-assisted Synthesis of Various C5-modified Pyrimidine Pyranonucleosides as Potential Cytotoxic Antitumor Agents. Current Microwave Chemistry, 2018, 4, .	0.8	1
118	A Proteomic Survey Indicates Sortilin as a Secondary Substrate of the ER Translocation Inhibitor Cyclotriazadisulfonamide (CADA). Molecular and Cellular Proteomics, 2017, 16, 157-167.	3.8	17
119	Structure-activity relationship studies on a Trp dendrimer with dual activities against HIV and enterovirus A71. Modifications on the amino acid. Antiviral Research, 2017, 139, 32-40.	4.1	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. Organic and Biomolecular Chemistry, 2017, 15, 1130-1139.	2.8	17
121	Design, Synthesis, and the Biological Evaluation of a New Series of Acyclic 1,2,3â€Triazole Nucleosides. Archiv Der Pharmazie, 2017, 350, 1700166.	4.1	8
122	Expanding the Antiviral Spectrum of 3-Fluoro-2-(phosphonomethoxy)propyl Acyclic Nucleoside Phosphonates: Diamyl Aspartate Amidate Prodrugs. Journal of Medicinal Chemistry, 2017, 60, 6220-6238.	6.4	22
123	Synthesis of different types of alkoxy fullerene derivatives from chlorofullerene C <sub>60</sub> Cl <sub>6</sub> . Organic and Biomolecular Chemistry, 2017, 15, 773-777.	2.8	28
124	Synthesis, anti-varicella-zoster virus and anti-cytomegalovirus activity of quinazoline-2,4-diones containing isoxazolidine and phosphonate substructures. European Journal of Medicinal Chemistry, 2017, 126, 84-100.	5.5	27
125	Signaling properties of the human chemokine receptors CXCR4 and CXCR7 by cellular electric impedance measurements. PLoS ONE, 2017, 12, e0185354.	2.5	21
126	Comparison of cell-based assays for the identification and evaluation of competitive CXCR4 inhibitors. PLoS ONE, 2017, 12, e0176057.	2.5	33

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127	Immunoregulatory effects of multipotent adult progenitor cells in a porcine ex vivo lung perfusion model. Stem Cell Research and Therapy, 2017, 8, 159.	5.5	51
128	Synthesis and Ativiral Activity of 5-(Benzylthio)-4-carbamyl-1,2,3-triazoles Against Human Cytomegalovirus (CMV) and Varicella-zoster Virus (VZV). Medicinal Chemistry, 2017, 13, 453-464.	1.5	9
129	A porcine ex vivo lung perfusion model with maximal argon exposure to attenuate ischemia-reperfusion injury. Medical Gas Research, 2017, 7, 28.	2.3	8
130	New Isoxazolidine-Conjugates of Quinazolinones—Synthesis, Antiviral and Cytostatic Activity. Molecules, 2016, 21, 959.	3.8	19
131	Design, Synthesis and Biological Evaluation of Novel Primaquine-Cinnamic Acid Conjugates of the Amide and Acylsemicarbazide Type. Molecules, 2016, 21, 1629.	3.8	22
132	Development and Identification of a Novel Anti-HIV-1 Peptide Derived by Modification of the N-Terminal Domain of HIV-1 Integrase. Frontiers in Microbiology, 2016, 7, 845.	3.5	13
133	Synthesis and the Biological Activity of Phosphonylated 1,2,3-Triazolenaphthalimide Conjugates. Molecules, 2016, 21, 1420.	3.8	12
134	Lectin-Like Molecules of Lactobacillus rhamnosus GG Inhibit Pathogenic Escherichia coli and Salmonella Biofilm Formation. PLoS ONE, 2016, 11, e0161337.	2.5	79
135	Membraneâ€permeable Triphosphate Prodrugs of Nucleoside Analogues. Angewandte Chemie - International Edition, 2016, 55, 5255-5258.	13.8	57
136	The lectin-like protein 1 in Lactobacillus rhamnosus GR-1 mediates tissue-specific adherence to vaginal epithelium and inhibits urogenital pathogens. Scientific Reports, 2016, 6, 37437.	3.3	38
137	Synthesis of novel N-acyl-β-d-glucopyranosylamines and ureas as potential lead cytostatic agents. Medicinal Chemistry Research, 2016, 25, 932-940.	2.4	1
138	Novel urea and bis -urea primaquine derivatives with hydroxyphenyl or halogenphenyl substituents: Synthesis and biological evaluation. European Journal of Medicinal Chemistry, 2016, 124, 622-636.	5.5	27
139	Steroids can reduce warm ischemic reperfusion injury in a porcine donation after circulatory death model with <i>ex vivo</i> lung perfusion evaluation. Transplant International, 2016, 29, 1237-1246.	1.6	42
140	Synthesis and antiviral properties of new derivatives of 2-(alkylsulfanyl)-6-[1-(2,6-difluorophenyl)cyclopropyl]-5-methylpyrimidin-4(3H)-one. Russian Journal of Organic Chemistry, 2016, 52, 1188-1193.	0.8	3
141	Lectin-Glycan Interaction Network-Based Identification of Host Receptors of Microbial Pathogenic Adhesins. MBio, 2016, 7, .	4.1	48
142	Design, Synthesis, and Antiviral Activity of Novel Ribonucleosides of 1,2,3â€Triazolylbenzylâ€aminophosphonates. Archiv Der Pharmazie, 2016, 349, 30-41.	4.1	30
143	Design, synthesis, and cytostatic activity of novel pyrazine sorafenib analogs. Medicinal Chemistry Research, 2016, 25, 2729-2741.	2.4	7
144	Humoral immunity in phenotypes of chronic lung allograft dysfunction: A broncho-alveolar lavage fluid analysis. Transplant Immunology, 2016, 38, 27-32.	1.2	36

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145	Aspartateâ€Based CXCR4 Chemokine Receptor Binding of Crossâ€Bridged Tetraazamacrocyclic Copper(II) and Zinc(II) Complexes. Chemistry - A European Journal, 2016, 22, 12916-12930.	3.3	16
146	High mannose-specific lectin Msl mediates key interactions of the vaginal Lactobacillus plantarum isolate CMPG5300. Scientific Reports, 2016, 6, 37339.	3.3	29
147	Novel isoxazolidine analogues of homonucleosides and homonucleotides. Tetrahedron, 2016, 72, 8294-8308.	1.9	9
148	Acyclic nucleoside phosphonates containing the amide bond. Monatshefte Für Chemie, 2016, 147, 2163-2177.	1.8	2
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