Konrad Misiura

List of Publications by Year in descending order

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567281 610901 48 652 15 24 citations h-index g-index papers 55 55 55 642 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Diastereomers of Nucleoside 3'-O-(2-Thio-1,3,2-oxathia(selena)phospholanes): Building Blocks for Stereocontrolled Synthesis of Oligo(nucleoside phosphorothioate)s. Journal of the American Chemical Society, 1995, 117, 12019-12029.	13.7	135
2	Discovery of tropinone-thiazole derivatives as potent caspase 3/7 activators, and noncompetitive tyrosinase inhibitors with high antiproliferative activity: Rational design, one-pot tricomponent synthesis, and lipophilicity determination. European Journal of Medicinal Chemistry, 2019, 175, 162-171.	5.5	37
3	Synthesis, antimicrobial evaluation and theoretical prediction of NMR chemical shifts of thiazole and selenazole derivatives with high antifungal activity against Candida spp Journal of Molecular Structure, 2016, 1108, 427-437.	3.6	31
4	Synthesis, antimicrobial and anticonvulsant screening of small library of tetrahydro-2H-thiopyran-4-yl based thiazoles and selenazoles. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 24-39.	5.2	28
5	Synthesis and anticonvulsant activities of novel 2-(cyclopentylmethylene)hydrazinyl-1,3-thiazoles in mouse models of seizures. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1576-1582.	5.2	25
6	Synthesis, In Vitro Biological Screening and Molecular Docking Studies of Novel Camphor-Based Thiazoles. Medicinal Chemistry, 2014, 10, 600-608.	1.5	24
7	Synthesis and In Vitro Antiproliferative Activity of Thiazole-Based Nitrogen Mustards: The Hydrogen Bonding Interaction between Model Systems and Nucleobases. Anti-Cancer Agents in Medicinal Chemistry, 2014, 14, 1271-1281.	1.7	22
8	Synthesis of Nucleoside α-Thiotriphosphates via an Oxathiaphospholane Approachâ€. Organic Letters, 2005, 7, 2217-2220.	4.6	21
9	Thiazole-based nitrogen mustards: Design, synthesis, spectroscopic studies, DFT calculation, molecular docking, and antiproliferative activity against selected human cancer cell lines. Journal of Molecular Structure, 2016, 1119, 139-150.	3.6	21
10	Synthesis and antitumor activity of analogs of ifosfamide modified in the N-(2-chloroethyl) group. Journal of Medicinal Chemistry, 1988, 31, 226-230.	6.4	19
11	The First Stereocontrolled Solid-Phase Synthesis of Di-, Tri-, and Tetra[adenosine (2â€~,5â€~) phosphorothioate]s. Journal of Organic Chemistry, 1998, 63, 7097-7100.	3.2	19
12	Anti-Sense Oligodeoxynucleoside Phosphorothioates Nonspecifically Inhibit Invasion of Red Blood Cells by Malaria Parasites. Biochemical and Biophysical Research Communications, 1996, 218, 930-933.	2.1	17
13	Role of GSTM1, GSTP1, and GSTT1 Gene Polymorphism in Ifosfamide Metabolism Affecting Neurotoxicity and Nephrotoxicity in Children. Journal of Pediatric Hematology/Oncology, 2005, 27, 582-589.	0.6	17
14	Dithymidylyl- $3\hat{a}\in^2$ -phosphorofluoridates: new synthesis and stability under solvolytic conditions. Journal of the Chemical Society Chemical Communications, 1995, , 613-614.	2.0	16
15	A New Method for Distinguishing between Enantiomers and Racemates and Assignment of Enantiomeric Purity by Means of Solid-State NMR. Examples from Oxazaphosphorinanes. Chemistry - A European Journal, 2002, 8, 5007-5011.	3.3	15
16	Tropinone-Derived Alkaloids as Potent Anticancer Agents: Synthesis, Tyrosinase Inhibition, Mechanism of Action, DFT Calculation, and Molecular Docking Studies. International Journal of Molecular Sciences, 2020, 21, 9050.	4.1	15
17	Synthesis, Antibacterial Activity, Interaction with Nucleobase and Molecular Docking Studies of 4-Formylbenzoic Acid Based Thiazoles. Medicinal Chemistry, 2016, 12, 553-562.	1.5	14
18	Synthesis and chemical and enzymatic reactivity of thymidine 3′-O- and 5′-O-phosphorofluoridothioates. Chemical Communications, 1998, , 515-516.	4.1	13

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19	DBU-assisted 1,3,2-oxathiaphospholane ring-opening condensation with selected O-, S-, N- and C-nucleophiles. Tetrahedron Letters, 2004, 45, 4301-4305.	1.4	13
20	Synthesis, Antimicrobial Activities and Molecular Docking Studies of Novel 6-Hydroxybenzofuran-3(2H)-one Based 2,4-Disubstituted 1,3- Thiazoles. Letters in Drug Design and Discovery, 2013, 10, 798-807.	0.7	12
21	Discovery and Evaluation of Efficient Selenazoles with High Antifungal Activity Against Candida spp Medicinal Chemistry, 2015, 11, 118-127.	1.5	12
22	Stereochemistry of phosphorus-nitrogen bond cleavage. First crystal and structural assignment in cyclic phosphoramidofluoridates. Journal of Organic Chemistry, 1985, 50, 1815-1818.	3.2	11
23	Synthesis, chemical and enzymatic reactivity, and toxicity of dithymidylyl-3′,5′-phosphorofluoridate and -phosphorothiofluoridate. Bioorganic and Medicinal Chemistry, 2001, 9, 1525-1532.	3.0	11
24	Deoxyxylothymidine 3′-O-phosphorothioates: Synthesis, stereochemistry and stereocontrolled incorporation into oligothymidylates. Bioorganic and Medicinal Chemistry Letters, 1997, 7, 2651-2656.	2.2	10
25	The synthesis of $5\hat{a}\in^2$ -O-DMT-thymidine $3\hat{a}\in^2$ -O-(2-THIO-1,3,2-oxaselenaphospholane) and its possible application in stereocontrolled synthesis of oligo(nucleoside phosphorothioate)s. Bioorganic and Medicinal Chemistry Letters, 1994, 4, 1037-1040.	2,2	8
26	Ifosfamide. Metabolic Studies, New Therapeutic Approaches and New Analogs. Mini-Reviews in Medicinal Chemistry, 2006, 6, 395-400.	2.4	8
27	Synthesis and antimicrobial activities of novel 6-(1,3-thiazol-4-yl)-1,3-benzoxazol-2(3H)-one derivatives. Heterocyclic Communications, 2014, 20, .	1.2	7
28	Studies on enzymatic hydrolysis of thymidin- $3\hat{a}\in^2$ -yl thymidin- $5\hat{a}\in^2$ -yl phosphorofluoridates and the corresponding phosphorothiofluoridates. Chemical Communications, 1999, , 2115-2116.	4.1	6
29	(S)-(â^')-Bromofosfamide (CBM-11): synthesis and antitumor activity and toxicity in mice. Anti-Cancer Drugs, 2001, 12, 453-458.	1.4	6
30	Synthesis and Structural Studies of SP and RP Diastereomers of Deoxyxylothymidyl- $3\hat{a}\in^2$ -O-acetylthymidyl ($3\hat{a}\in^2$, $5\hat{a}\in^2$)-O-(2-Cyanoethyl)phosphorothioate in Solution and in the Solid State. European Journal of Organic Chemistry, 2001, 2001, 1491-1501.	2.4	5
31	Reactivity of nucleoside 5?-O-phosphates, -phosphorothioates, -methanephosphonates, and -methanephosphonothioates toward activated xylonucleosides. Heteroatom Chemistry, 1999, 10, 91-104.	0.7	4
32	Synthesis, in vitro metabolic studies, and antitumour activity of methyl analogues of ifosfamide. Archiv Der Pharmazie, 2001, 334, 291.	4.1	4
33	Studies on the Side-chain Hydroxylation of Ifosfamide and Its Bromo Analogue. Bioorganic and Medicinal Chemistry Letters, 2002, 12, 427-431.	2.2	4
34	Synthesis and biological evaluation of novel 2-(1H-imidazol-2-ylmethylidene)hydrazinyl- 1,3-thiazoles as potential antimicrobial agents. Heterocyclic Communications, 2015, 21, .	1.2	4
35	Synthesis and reactivity of dithymidylyl-3',5''-phosphorothiofluoridates. Collection of Czechoslovak Chemical Communications, 1996, 61, 101-106.	1.0	4
36	Isophosphoramide mustard analogues as prodrugs for anticancer gene-directed enzyme-prodrug therapy (GDEPT) Acta Biochimica Polonica, 2002, 49, 169-176.	0.5	4

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37	Synthesis and Antimicrobial Activities of (4,5,6,7-Tetrahydro-1H-indazol- 2(3H)-yl)thiazole Derivatives. Letters in Drug Design and Discovery, 2014, 11, 960-967.	0.7	4
38	Synthesis of 17O (and 18O) labelled isophosphoramide mustard. Journal of Labelled Compounds and Radiopharmaceuticals, 1994, 34, 247-254.	1.0	3
39	The synthesis, X-ray and solid state NMR studies of 2-N,N-diisopropylamino-1,3,2λ5-oxaselenaphospholane-2-selone. Journal of the Chemical Society Perkin Transactions II, 1997, , 163-168.	0.9	3
40	Synthesis and antitumour activity of stereoisomers of 4-hydroperoxy derivatives of ifosfamide and its bromo analogue. Il Farmaco, 2002, 57, 315-319.	0.9	3
41	Nucleophilic N ¹ → N ³ Rearrangement of 5′-O-Trityl-O ² ,3′-Cycloanhydrothymidine. Nucleosides, Nucleotides and Nucleic Acids, 2000, 19, 1657-1673.	1.1	2
42	Analysis of the Urinary Excretion of Ifosfamide and its N-Dechloroethylated Metabolites in Children Using 31P-NMR Spectroscopy. Arzneimittelforschung, 2003, 53, 372-377.	0.4	2
43	Triazene salts: Design, synthesis, ctDNA interaction, lipophilicity determination, DFT calculation, and antiproliferative activity against human cancer cell lines. Saudi Pharmaceutical Journal, 2019, 27, 303-311.	2.7	2
44	The Disulfide Analogues of Isophosphoramide Mustard for Anticancer Therapy. Letters in Drug Design and Discovery, 2014, 12, 172-179.	0.7	2
45	lon Exchange HPLC Analysis of Oligoribonucleotides and Chimeric Oligoribo-oligodeoxyribonucleotidesa. Annals of the New York Academy of Sciences, 1992, 660, 321-323.	3.8	1
46	Oxathiaphospholane Approach to the Synthesis of Nucleoside MethaneÂphosphonothioates. Synlett, 2004, 2004, 2143-2146.	1.8	0
47	Isophosphoramide mustard analogues as prodrugs for anticancer gene-directed enzyme-prodrug therapy (GDEPT). Acta Biochimica Polonica, 2002, 49, 169-76.	0.5	0
48	Phosphate prodrugs of isophosphoramide mustard. Acta Poloniae Pharmaceutica, 2003, 60, 109-12.	0.1	0