

Wolfgang Holzer

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1711001/publications.pdf>

Version: 2024-02-01

217
papers

4,546
citations

126708

33
h-index

174990

52
g-index

263
all docs

263
docs citations

263
times ranked

3708
citing authors

#	ARTICLE	IF	CITATIONS
1	Synthesis, Cytotoxicity, and Antitumor Activity of Copper(II) and Iron(II) Complexes of 4N-Azabicyclo[3.2.2]nonane Thiosemicarbazones Derived from Acyl Diazines. <i>Journal of Medicinal Chemistry</i> , 2001, 44, 2164-2171.	2.9	233
2	Increasing the Reactivity of Amides towards Organometallic Reagents: An Overview. <i>Advanced Synthesis and Catalysis</i> , 2014, 356, 3697-3736.	2.1	207
3	Exploiting a "Beast" in Carbenoid Chemistry: Development of a Straightforward Direct Nucleophilic Fluoromethylation Strategy. <i>Journal of the American Chemical Society</i> , 2017, 139, 13648-13651.	6.6	104
4	Structures of Highly Twisted Amides Relevant to Amide N=C Cross-Coupling: Evidence for Ground-State Amide Destabilization. <i>Chemistry - A European Journal</i> , 2016, 22, 14494-14498.	1.7	94
5	Addition of lithium carbenoids to isocyanates: a direct access to synthetically useful N-substituted 2-haloacetamides. <i>Chemical Communications</i> , 2013, 49, 8383.	2.2	85
6	Substituted 4-Acylpyrazoles and 4-Acylpyrazolones: Synthesis and Multidrug Resistance-Modulating Activity. <i>Journal of Medicinal Chemistry</i> , 1998, 41, 4001-4011.	2.9	83
7	Synthesis of pyrazole-based hybrid molecules: Search for potent multidrug resistance modulators. <i>Bioorganic and Medicinal Chemistry</i> , 2006, 14, 5061-5071.	1.4	82
8	Chemoselective Activation Strategies of Amidic Carbonyls towards Nucleophilic Reagents. <i>Australian Journal of Chemistry</i> , 2013, 66, 507.	0.5	78
9	Efficient Access to All-Carbon Quaternary and Tertiary α -Functionalized Homoallyl-type Aldehydes from Ketones. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 12677-12682.	7.2	71
10	Synthesis and evaluation of indole, pyrazole, chromone and pyrimidine based conjugates for tumor growth inhibitory activities – Development of highly efficacious cytotoxic agents. <i>European Journal of Medicinal Chemistry</i> , 2010, 45, 4968-4982.	2.6	67
11	Modular and Chemoselective Strategy for the Direct Access to α -Fluoroepoxides and Aziridines via the Addition of Fluoroiodomethylithium to Carbonyl-Like Compounds. <i>Organic Letters</i> , 2019, 21, 584-588.	2.4	65
12	Telescoped, Divergent, Chemoselective C1 and C1-C1 Homologation of Imine Surrogates: Access to Quaternary Chloro- and Halomethyl- and Trifluoromethyl Aziridines. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 2479-2484.	7.2	64
13	Pyridazines. 63. Novel thiosemicarbazones derived from formyl- and acyldiazines: synthesis, effects on cell proliferation, and synergism with antiviral agents. <i>Journal of Medicinal Chemistry</i> , 1992, 35, 3288-3296.	2.9	59
14	Bromomethylithium-mediated chemoselective homologation of disulfides to dithioacetals. <i>Chemical Communications</i> , 2016, 52, 2639-2642.	2.2	59
15	An easy access to anomeric glycosyl amides and imines (Schiff bases) via transformation of glycopyranosyl trimethylphosphinimides. <i>Tetrahedron</i> , 2001, 57, 4609-4621.	1.0	57
16	Synthesis of α,β -Unsaturated α -Haloketones through the Chemoselective Addition of Halomethylithiums to Weinreb Amides. <i>Journal of Organic Chemistry</i> , 2013, 78, 7764-7770.	1.7	57
17	Lithium Halomethylcarbenoids: Preparation and Use in the Homologation of Carbon Electrophiles. <i>Chemical Record</i> , 2016, 16, 2061-2076.	2.9	55
18	Identification of Ligand-Binding Regions of P-Glycoprotein by Activated-Pharmacophore Photoaffinity Labeling and Matrix-Assisted Laser Desorption/Ionization Time-of-Flight Mass Spectrometry. <i>Molecular Pharmacology</i> , 2002, 61, 637-648.	1.0	53

#	ARTICLE	IF	CITATIONS
19	Direct and Chemoselective Synthesis of Tertiary Difluoroketones via Weinreb Amide Homologation with a CHF ₂ -Carbene Equivalent. <i>Organic Letters</i> , 2019, 21, 8261-8265.	2.4	53
20	Evidence and isolation of tetrahedral intermediates formed upon the addition of lithium carbenoids to Weinreb amides and N-acylpyrroles. <i>Chemical Communications</i> , 2017, 53, 9498-9501.	2.2	52
21	4-Acyl-5-methyl-2-phenylpyrazolones: NMR and X-Ray Structure Investigations. <i>Heterocycles</i> , 1999, 50, 799.	0.4	49
22	On the tautomerism of pyrazolones: the geminal 2J[pyrazole C-4,H-3(5)] spin coupling constant as a diagnostic tool. <i>Tetrahedron</i> , 2004, 60, 6791-6805.	1.0	49
23	Chemoselective Addition of Halomethylolithiums to Functionalized Isatins: A Straightforward Access to Spiro-Epoxyindoles. <i>Advanced Synthesis and Catalysis</i> , 2016, 358, 172-177.	2.1	47
24	Pd-catalyzed cross-coupling reactions of halogenated 1-phenylpyrazol-3-ols and related triflates. <i>Tetrahedron</i> , 2009, 65, 7817-7824.	1.0	45
25	Homologation of Isocyanates with Lithium Carbenoids: A Straightforward Access to α -Halomethyl- and α,α -Dihalomethylamides. <i>Synthesis</i> , 2014, 46, 2897-2909.	1.2	45
26	Highly efficient synthesis of functionalized α -oxyketones via Weinreb amides homologation with α -oxygenated organolithiums. <i>Chemical Communications</i> , 2016, 52, 7584-7587.	2.2	44
27	Compounds from <i>Caesalpinia sappan</i> with anti-inflammatory properties in macrophages and chondrocytes. <i>Food and Function</i> , 2016, 7, 1671-1679.	2.1	44
28	Chemoselective Synthesis of α -Substituted α -Amino- α,α -Chloro Ketones via Chloromethylation of Glycine-Derived Weinreb Amides. <i>Advanced Synthesis and Catalysis</i> , 2013, 355, 919-926.	2.1	41
29	Chemoselective efficient synthesis of functionalized α -oxonitriles through cyanomethylation of Weinreb amides. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 1969-1973.	1.5	41
30	Expeditious and Chemoselective Synthesis of α -Aryl and α -Alkyl Selenomethylketones via Homologation Chemistry. <i>Organic Letters</i> , 2018, 20, 2685-2688.	2.4	39
31	A Robust, Eco-Friendly Access to Secondary Thioamides through the Addition of Organolithium Reagents to Isothiocyanates in Cyclopentyl Methyl Ether (CPME). <i>Chemistry - A European Journal</i> , 2015, 21, 18966-18970.	1.7	38
32	Highly efficient chemoselective N-TBS protection of anilines under exceptional mild conditions in the eco-friendly solvent 2-methyltetrahydrofuran. <i>Green Chemistry</i> , 2011, 13, 1986.	4.6	37
33	Heterocyclic Analogs of Thioflavones: Synthesis and NMR Spectroscopic Investigations. <i>Molecules</i> , 2009, 14, 3814-3832.	1.7	34
34	Spiro-Fused (C2)-Azirino-(C4)-pyrazolones, a New Heterocyclic System. Synthesis, Spectroscopic Studies and X-ray Structure Analysis. <i>Journal of Organic Chemistry</i> , 2003, 68, 7943-7950.	1.7	32
35	Direct and Chemoselective Electrophilic Monofluoromethylation of Heteroatoms (O-, S-, N-). <i>Tetrahedron Letters</i> , 2014, 55, 1078-1081.	2.4	32
36	Synthesis of 3-substituted 1-phenyl-1H-pyrazole-4-carbaldehydes and the corresponding ethanones by Pd-catalysed cross-coupling reactions. <i>Arkivoc</i> , 2011, 2011, 1-21.	0.3	31

#	ARTICLE	IF	CITATIONS
37	Synthesis of substituted 3-phenyl-4-pyrazolo[4,3-d]isoxazoles from corresponding 4-benzoyl-5-hydroxypyrazoles. <i>Journal of Heterocyclic Chemistry</i> , 2003, 40, 303-308.	1.4	30
38	Robust eco-friendly protocol for the preparation of β -hydroxy- α , β -acetylenic esters by sequential one-pot elimination-addition of 2-bromoacrylates to aldehydes promoted by LTMP in 2-MeTHF. <i>Green Chemistry</i> , 2012, 14, 1859.	4.6	30
39	Chemoselective Additions of Chloromethylithium Carbenoid to Cyclic Enones: A Direct Access to Chloromethyl Allylic Alcohols. <i>Advanced Synthesis and Catalysis</i> , 2014, 356, 1761-1766.	2.1	30
40	First Synthesis of 3-Acetyl-2-aminothiophenes Using the Gewald Reaction. <i>Molecules</i> , 2006, 11, 371-376.	1.7	29
41	Pyrazolo[4 α ,3 β :5,6]pyrano[2,3 α :b β]quinoxalin-4(1H)-one: Synthesis and characterization of a novel tetracyclic ring system. <i>Journal of Heterocyclic Chemistry</i> , 2007, 44, 1139-1143.	1.4	29
42	A study in desmotropy. <i>Solid State Nuclear Magnetic Resonance</i> , 2008, 34, 68-76.	1.5	29
43	<i>Eulophia macrobulbon</i> - an orchid with significant anti-inflammatory and antioxidant effect and anticancerogenic potential exerted by its root extract. <i>Phytomedicine</i> , 2017, 24, 157-165.	2.3	29
44	Pyridazines 47. The Configuration of Novel Thiosemicarbozone Derivatives of Pyridazinecarbaldehydes and Alkyl Pyridazinyl Ketones. <i>Heterocycles</i> , 1989, 29, 1399.	0.4	28
45	Convenient and rapid determination of the configuration of aldoximes and ketoximes by means of noe difference spectroscopy. <i>Tetrahedron Letters</i> , 1990, 31, 3109-3112.	0.7	28
46	The [2-(Trimethylsilyl)ethoxy]methyl Function as a Suitable N-1 Protecting Group in Lithiation Reactions with Pyrazoles and 1,2,4-Triazoles. <i>Heterocycles</i> , 1992, 34, 303.	0.4	28
47	Synthesis and characterization of 4,5-dihydro-1H-pyrazolo[3,4b][1,4]azaphosphinines. <i>Heteroatom Chemistry</i> , 1999, 10, 391-398.	0.4	27
48	Acridone based Cu ²⁺ -responsive ON/OFF key pad. <i>Sensors and Actuators B: Chemical</i> , 2010, 150, 50-56.	4.0	26
49	<i>Cajanus cajan</i> - a source of PPAR β activators leading to anti-inflammatory and cytotoxic effects. <i>Food and Function</i> , 2016, 7, 3798-3806.	2.1	26
50	NOE difference spectroscopy as a versatile tool for spectral and structural assignment in various N-1 substituted pyrazoles. <i>Tetrahedron</i> , 1991, 47, 1393-1398.	1.0	25
51	The 4-Methoxybenzyl (PMB) Function as a Versatile Protecting Group in the Synthesis of N-Unsubstituted Pyrazolones. <i>Heterocycles</i> , 2004, 63, 2537.	0.4	25
52	Synthesis and ring transformations of 1-amino-1,2,3,9a-tetrahydroimidazo[1,2-a]indol-2(9H)-ones. <i>Tetrahedron</i> , 2006, 62, 3309-3319.	1.0	25
53	Chemoselective reduction of isothiocyanates to thioformamides mediated by the Schwartz reagent. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 1970-1978.	1.5	25
54	Pseudo-Dipeptide Bearing β , β -Difluoromethyl Ketone Moiety as Electrophilic Warhead with Activity against Coronaviruses. <i>International Journal of Molecular Sciences</i> , 2021, 22, 1398.	1.8	25

#	ARTICLE	IF	CITATIONS
55	On the application of NOE difference spectroscopy for spectral and structural assignments with substituted 1,2,3-triazoles. <i>Tetrahedron</i> , 1991, 47, 9783-9792.	1.0	24
56	Highly chemoselective difluoromethylative homologation of iso(thio)cyanates: expeditious access to unprecedented 1,1-difluoro(thio)amides. <i>Chemical Communications</i> , 2019, 55, 12960-12963.	2.2	24
57	Alkylation of Pyrazolones via the Mitsunobu Reaction. <i>Heterocycles</i> , 1997, 45, 309.	0.4	23
58	Efficient Access to All-Carbon Quaternary and Tertiary α -Functionalized Homoallyl-type Aldehydes from Ketones. <i>Angewandte Chemie</i> , 2017, 129, 12851-12856.	1.6	23
59	Sustainable Asymmetric Organolithium Chemistry: Enantio- and Chemoselective Acylations through Recycling of Solvent, Sparteine, and Weinreb α -Amine. <i>ChemSusChem</i> , 2019, 12, 1147-1154.	3.6	23
60	Chemoselective Homologation-Deoxygenation Strategy Enabling the Direct Conversion of Carbonyls into ($n+1$)-Halomethyl-Alkanes. <i>Organic Letters</i> , 2020, 22, 7629-7634.	2.4	23
61	Thiophen als Strukturelement physiologisch aktiver Substanzen, 12. Mitt. Thiophenanaloga antiviraler Chalkone. <i>Archiv Der Pharmazie</i> , 1985, 318, 48-59.	2.1	22
62	N-Substituted ethyl 4-pyrazolecarboxylates: Synthesis and spectroscopic investigations. <i>Journal of Heterocyclic Chemistry</i> , 1993, 30, 865-872.	1.4	22
63	Tri- and Tetracyclic Heteroaromatic Systems: Synthesis of Novel Benzo-, Benzothieno- and Thieno-Fused Pyrano[2,3-c]pyrazol-4(1H)-ones. <i>Heterocycles</i> , 2007, 71, 87.	0.4	22
64	Synthesis and biological evaluation of novel cytotoxic azanaphthoquinone annelated pyrrolo oximes. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 6091-6095.	1.0	22
65	Sonogashira-type Reactions with 5-Chlorophenyl-1H-pyrazole-4-carbaldehydes: A Straightforward Approach to Pyrazolo[4,3-f]pyridines. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 5123-5133.	1.2	22
66	Highly efficient and environmentally benign preparation of Weinreb amides in the biphasic system 2-MeTHF/water. <i>RSC Advances</i> , 2013, 3, 10158.	1.7	22
67	Metal-Free Intramolecular Alkyne-Azide Cycloaddition To Construct the Pyrazolo[4,3-f][1,2,3]triazolo[5,1-c][1,4]oxazepine Ring System. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 5663-5670.	1.2	22
68	Eco-friendly chemoselective N-functionalization of isatins mediated by supported KF in 2-MeTHF. <i>Green Chemistry</i> , 2015, 17, 4194-4197.	4.6	22
69	New 1-Substituted 4-Cinnamoyl-5-hydroxypyrazoles and Precursors thereof: Synthesis, Ring Closure Reactions and NMR-Spectroscopic Investigations. <i>Heterocycles</i> , 2003, 60, 2323.	0.4	22
70	Ethyl- and Triflyloxy-1H-pyrazole-4-carboxylates in the Synthesis of Condensed Pyrazoles by Pd-Catalysed Cross-Coupling Reactions. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 1880-1890.	1.2	21
71	A greener and efficient access to substituted four- and six-membered sulfur-bearing heterocycles. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 5000-5015.	1.5	21
72	Merging lithium carbenoid homologation and enzymatic reduction: A combinative approach to the HIV-protease inhibitor Nelfinavir. <i>Tetrahedron</i> , 2018, 74, 2211-2217.	1.0	21

#	ARTICLE	IF	CITATIONS
73	Synthesis and NMR Spectroscopic Investigations with 3-Amino-, 3-Hydroxy-, and 3-Methoxy-4-acyl-1-phenyl-2-pyrazolin-5-ones. <i>Heterocycles</i> , 2004, 63, 1311.	0.4	20
74	A Convenient Approach to Heterocyclic Building Blocks: Synthesis of Novel Ring Systems Containing a [5,6]Pyrano[2,3-c]pyrazol-4(1H)-one Moiety. <i>Molecules</i> , 2007, 12, 60-73.	1.7	20
75	Highly chemoselective synthesis of aryl allylic sulfoxides through calcium hypobromite oxidation of aryl allylic sulfides. <i>Tetrahedron Letters</i> , 2012, 53, 967-972.	0.7	20
76	Consecutive and Selective Double Methylene Insertion of Lithium Carbenoids to Isothiocyanates: A Direct Assembly of Four-Membered Sulfur-Containing Cycles. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 24854-24858.	7.2	20
77	Pyrazoles 3. N-1 Protected 4-Substituted Pyrazoles " Synthesis and Nmr Investigation. <i>Heterocycles</i> , 1988, 27, 2443.	0.4	19
78	On the discrimination of tetrazole regioisomers by NOE difference spectroscopy. <i>Monatshefte für Chemie</i> , 1992, 123, 1027-1036.	0.9	19
79	Functionalisation of 1,2,3-triazole via lithiation of 1-((trimethylsilyl)ethoxy)methyl-1,2,3-triazole. <i>Journal of Heterocyclic Chemistry</i> , 1992, 29, 1203-1207.	1.4	19
80	Heterocyclic Analogues of Xanthone and Xanthione. 1H-Pyrano[2,3-c:6,5-c]dipyrazol-4(7H)-ones and Thiones: Synthesis and NMR Data. <i>Molecules</i> , 2010, 15, 6106-6126.	1.7	19
81	Substituted Sulfur Methyl Carbanions: Effective Homologating Agents for the Chemoselective Preparation of Oxo Thioethers from Weinreb Amides. <i>European Journal of Organic Chemistry</i> , 2018, 2018, 2466-2470.	1.2	19
82	Pyrazoles. 5. Novel pyrazole analogues of flavanone, flavone and flavane. <i>Journal of Heterocyclic Chemistry</i> , 1991, 28, 1047-1050.	1.4	18
83	Spectral and structural assignments with various N-substituted 1,2,4-triazoles: Noe difference spectroscopy as a powerful tool. <i>Tetrahedron</i> , 1991, 47, 5471-5480.	1.0	18
84	Configurational assignments of oximes derived from 5-formyl and 5-cyano-1,2,4-triazines. <i>Journal of Heterocyclic Chemistry</i> , 1993, 30, 413-418.	1.4	18
85	The Structure of 4-Benzoyl-5-methyl-2-phenylpyrazol-3-one Oxime and Its Methyl Derivatives. <i>European Journal of Organic Chemistry</i> , 2003, 2003, 1209-1219.	1.2	18
86	Synthesis of electroactive hydrazones derived from 3-(10-alkyl-10H-phenothiazin-3-yl)-2-propenals and their corresponding 3,3-bispropenals. <i>Tetrahedron</i> , 2012, 68, 3552-3559.	1.0	18
87	Chemoselective Ca-Mediated Acylation of Alcohols and Amines in 2-Methyltetrahydrofuran. <i>ChemSusChem</i> , 2013, 6, 905-910.	3.6	18
88	Homologation of halostannanes with carbenoids: a convenient and straightforward one-step access to α -functionalized organotin reagents. <i>Chemical Communications</i> , 2018, 54, 10112-10115.	2.2	18
89	Pyrazoles. 6. Synthesis of novel heteroaryl 4-pyrazolyl ketones. <i>Journal of Heterocyclic Chemistry</i> , 1991, 28, 1189-1192.	1.4	17
90	Structure/Odor Relationships of (-)- and (+)- β -Vetivone, and Their Demethyl Derivatives. <i>Helvetica Chimica Acta</i> , 1998, 81, 2292-2299.	1.0	17

#	ARTICLE	IF	CITATIONS
91	An Efficient Approach to Heterocyclic Analogues of Xanthone: A Short Synthesis of All Possible Pyrido[5,6]pyrano[2,3-c]pyrazol-4(1H)-ones. <i>Synthesis</i> , 2006, 2006, 4219-4229.	1.2	17
92	Novel fluoro-substituted benzo- and benzothieno fused pyrano[2,3-c]pyrazol-4(1H)-ones. <i>Journal of Fluorine Chemistry</i> , 2010, 131, 1013-1024.	0.9	17
93	On the Tautomerism of N-Substituted Pyrazolones: 1,2-Dihydro-3H-pyrazol-3-ones versus 1H-Pyrazol-3-ols. <i>Molecules</i> , 2018, 23, 129.	1.7	17
94	Halogen-impacted Reactivity in Lithium Carbenoid Mediated Homologations of Imine Surrogates: Direct Assembly of bis-trifluoromethyl-diketiminates and the Dual Role of LiCH ₂ I. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 20852-20857.	7.2	17
95	On the application of homonuclear NOE difference spectroscopy as a convenient tool for configurational assignment of compounds with a C=N bond. <i>Monatshefte für Chemie</i> , 1990, 121, 837-846.	0.9	16
96	NMR spectroscopic investigations with ethyl (hetero)aryl-5-hydroxy-1H-pyrazole-4-carboxylates. <i>Journal of Heterocyclic Chemistry</i> , 1995, 32, 1341-1349.	1.4	16
97	4-Acyl-5-hydroxy-1-phenyl-3-trifluoromethylpyrazoles: Synthesis and NMR Spectral Investigations. <i>Heterocycles</i> , 2006, 68, 1825.	0.4	16
98	Highly efficient and chemoselective α -iodination of acrylate esters through Morita-Baylis-Hillman-type chemistry. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 1085.	1.5	16
99	Use of activated enol ethers in the synthesis of pyrazoles: reactions with hydrazine and a study of pyrazole tautomerism. <i>Beilstein Journal of Organic Chemistry</i> , 2014, 10, 752-760.	1.3	16
100	On the Bioisosteric Potential of Diazines: Diazine Analogues of the Combined Thromboxane A ₂ Receptor Antagonist and Synthetase Inhibitor Ridogrel. <i>Journal of Medicinal Chemistry</i> , 1996, 39, 4058-4064.	2.9	15
101	Synthesis and NMR-Investigation of Annelated Pyrrole Derivatives. <i>Heterocycles</i> , 1997, 45, 1989.	0.4	15
102	Sonogashira Coupling Offers a New Synthetic Route to Thieno[2,3-c]pyrazoles. <i>Synthetic Communications</i> , 2011, 41, 541-547.	1.1	15
103	Synthesis and anti-mitotic activity of 2,4- or 2,6-disubstituted- and 2,4,6-trisubstituted-2H-pyrazolo[4,3-c]pyridines. <i>European Journal of Medicinal Chemistry</i> , 2018, 150, 908-919.	2.6	15
104	Pyridazines. XXVI. A novel synthesis of pyrano[2,3-d]pyridazines. <i>Journal of Heterocyclic Chemistry</i> , 1986, 23, 93-96.	1.4	14
105	¹³ C nuclear magnetic resonance spectra of 3,6-disubstituted pyridazines. <i>Canadian Journal of Chemistry</i> , 1991, 69, 972-977.	0.6	14
106	On the structure of guanylhya zones derived from aromatic aldehydes. <i>Monatshefte für Chemie</i> , 1992, 123, 1163-1173.	0.9	14
107	Synthesis and ¹³ C NMR study of some N-substituted 4-iodo- and 3,4-diiodopyrazoles. <i>Journal of Heterocyclic Chemistry</i> , 1995, 32, 189-194.	1.4	14
108	Synthesis and Odor of Chiral Partial Structures of Khusimone. Part 1. <i>Helvetica Chimica Acta</i> , 1997, 80, 139-145.	1.0	14

#	ARTICLE	IF	CITATIONS
109	2, 3-Diaryl-5-ethylsulfanylmethyltetrahydrofurans as a new class of COX-2 inhibitors and cytotoxic agents. <i>Organic and Biomolecular Chemistry</i> , 2008, 6, 2706.	1.5	14
110	On the Tautomerism of Cinnolin-4-ol, Cinnolin-4-thiol, and Cinnolin-4-amine. <i>Heterocycles</i> , 2008, 75, 77.	0.4	14
111	Synthesis of pyrazolo[4- <i>b</i> ,3- <i>e</i> :3,4]pyrido[1,2- <i>a</i>]benzimidazoles and related new ring systems by tandem cyclisation of vic-alkynylpyrazole-4-carbaldehydes with (het)aryl-1,2-diamines and investigation of their optical properties. <i>Tetrahedron</i> , 2015, 71, 3385-3395.	1.0	14
112	Straightforward chemoselective access to unsymmetrical dithioacetals through a thiosulfonate homologation-nucleophilic substitution sequence. <i>Chemical Communications</i> , 2020, 56, 12395-12398.	2.2	14
113	<i>N</i> -substituted 3,5-dimethoxy-4-halogeno-1 <i>H</i> -pyrazoles: Synthesis and NMR study. <i>Journal of Heterocyclic Chemistry</i> , 1995, 32, 1351-1354.	1.4	13
114	Aryl Diazinyl Ketoximes: Synthesis and Configurational Assignment. <i>Heterocycles</i> , 1996, 43, 151.	0.4	13
115	Synthesis of trifluoromethyl-substituted pyrazolo[4,3- <i>c</i>]pyridines – sequential versus multicomponent reaction approach. <i>Beilstein Journal of Organic Chemistry</i> , 2014, 10, 1759-1764.	1.3	13
116	Molecular dimensions and structural features of neutral polysaccharides from the seed mucilage of <i>Hyptis suaveolens</i> L.. <i>Food Chemistry</i> , 2017, 221, 1997-2004.	4.2	13
117	±-Arylamino Diazoketones: Diazomethane-Loading Controlled Synthesis, Spectroscopic Investigations, and Structural X-ray Analysis. <i>Journal of Organic Chemistry</i> , 2018, 83, 4336-4347.	1.7	13
118	Synthesis and anthelmintic activity of benzopyrano[2,3- <i>c</i>]pyrazol-4(2H)-one derivatives. <i>Molecular Diversity</i> , 2020, 24, 1025-1042.	2.1	13
119	Electrophilicity Scale of Activated Amides: 17 ¹⁷ O and 15 ¹⁵ N NMR Chemical Shifts of Acyclic Twisted Amides in ¹³ C(O) Cross-Coupling. <i>Chemistry - A European Journal</i> , 2020, 26, 16246-16250.	1.7	13
120	Beiträge zur Chemie des Pyrazolsystems, 1. Mitt.: Ein effizienter Zugang zu Aryl- oder Benzyl- ⁴ -pyrazolylketonen und ⁴ -carbinolen. <i>Archiv Der Pharmazie</i> , 1987, 320, 1267-1272.	2.1	12
121	NMR spectroscopic investigations with isatin guanylhydrazones. <i>Journal of Heterocyclic Chemistry</i> , 1996, 33, 675-680.	1.4	12
122	Synthesis and biological evaluation of new cytotoxic azanaphthoquinone pyrrolo-annelated derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 3950-3952.	1.0	12
123	¹⁷ O NMR and ¹⁵ N NMR chemical shifts of sterically-hindered amides: ground-state destabilization in amide electrophilicity. <i>Chemical Communications</i> , 2019, 55, 4423-4426.	2.2	12
124	Highly Regioselective and Efficient Synthesis of Aminoepoxides by Ring Closure of Aminohalohydrins Mediated by KF-Celite. <i>Synlett</i> , 2011, 2011, 1831-1834.	1.0	11
125	Synthesis of stable ±-fluoromethyl putative carbanions via a chemoselective reduction-monofluoromethylation sequence of diselenides under sustainable conditions. <i>Tetrahedron</i> , 2021, 85, 131921.	1.0	11
126	Synthesis and Odor of Chiral Partial Structures of Khusimone. Part 3. Short communication. <i>Helvetica Chimica Acta</i> , 1998, 81, 40-45.	1.0	10

#	ARTICLE	IF	CITATIONS
127	Synthesis of anticancer compounds, III (Bioorg Med Chem Lett 17, 6091, 2007), carbinol derivatives of azanaphthoquinone annelated pyrroles. Monatshefte für Chemie, 2009, 140, 309-313.	0.9	10
128	A straightforward and general access to α -phthalimido- β -substituted propan-2-ones. Tetrahedron Letters, 2012, 53, 5106-5109.	0.7	10
129	Multinuclear NMR spectra and GIAO/DFT calculations of N-benzylazoles and N-benzylbenzazoles. Structural Chemistry, 2019, 30, 1729-1735.	1.0	10
130	Taking advantage of lithium monohalocarbenoid intrinsic β -elimination in 2-MeTHF: controlled epoxide ring-opening en route to halohydrins. Organic and Biomolecular Chemistry, 2021, 19, 2038-2043.	1.5	10
131	Pyridazines. LI. On the Reactivity of Pyridazine-carbaldehydes towards Selected Active-Hydrogen Compounds. Journal of Heterocyclic Chemistry, 1990, 27, 1313-1321.	1.4	9
132	Determination of the Stereochemistry of Chemotherapeutics Derived from 5-Nitrofurfural: NOE Difference Spectroscopy as a Simple and Reliable Method. Archiv Der Pharmazie, 1992, 325, 769-772.	2.1	9
133	Synthesis and Odor of Chiral Partial Structures of Khusimone. Part 2. Helvetica Chimica Acta, 1997, 80, 1857-1864.	1.0	9
134	Synthesis of Azanaphthoquinone Annelated Pyrroles. Heterocycles, 2001, 54, 111.	0.4	9
135	Derivatives of pyrazinecarboxylic acid: ^1H , ^{13}C and ^{15}N NMR spectroscopic investigations. Magnetic Resonance in Chemistry, 2009, 47, 617-624.	1.1	9
136	Dipyrazolo[1,5-a:4',3'-c]pyridines – a new heterocyclic system accessed via multicomponent reaction. Beilstein Journal of Organic Chemistry, 2012, 8, 2223-2229.	1.3	9
137	Chemoselective oxidative hydrolysis of EWG protected α -arylamino vinyl bromides to α -arylamino- β -bromoacetones. Tetrahedron Letters, 2013, 54, 4369-4372.	0.7	9
138	Development of potential selective and reversible pyrazoline based MAO-B inhibitors as MAO-B PET tracer precursors and reference substances for the early detection of Alzheimer's disease. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 4490-4495.	1.0	9
139	Ring-closing metathesis as a key step to construct 2,6-dihydropyrano[2,3-c]pyrazole ring system. Arkivoc, 2018, 2018, 296-307.	0.3	9
140	A Straightforward Homologation of Carbon Dioxide with Magnesium Carbenoids en Route to α -Halocarboxylic Acids. Advanced Synthesis and Catalysis, 2019, 361, 1001-1006.	2.1	9
141	Design, Synthesis, and Pharmacological Evaluation of Novel α -Subunit-Selective β -Aminobutyric Acid Type A (GABA _A) Receptor Modulators. Journal of Medicinal Chemistry, 2019, 62, 317-341.	2.9	9
142	Configurational assignment of aryl heteroaryl ketoximes by means of homonuclear NOE difference spectroscopy. Collection of Czechoslovak Chemical Communications, 1991, 56, 2251-2257.	1.0	9
143	N-Substituted Bromopyrazoles: Synthesis and ^{13}C Nmr Study. Heterocycles, 1994, 38, 2433.	0.4	8
144	Cyclization reactions of N1-(glycopyranosylamino) guanidines. Carbohydrate Research, 1997, 302, 229-235.	1.1	8

#	ARTICLE	IF	CITATIONS
145	Synthesis of in vivo Metabolites of the New Adenosine A3 Receptor PET-Radiotracer [18F]FE@SUPPY. Heterocycles, 2008, 75, 339.	0.4	8
146	Synthesis of 2 <i>H</i> -furo[2,3- <i>c</i>]pyrazole ring systems through silver(I) ion-mediated ring-closure reaction. Beilstein Journal of Organic Chemistry, 2019, 15, 679-684.	1.3	8
147	Enhanced arecoline derivatives as muscarinic acetylcholine receptor M1 ligands for potential application as PET radiotracers. European Journal of Medicinal Chemistry, 2020, 204, 112623.	2.6	8
148	Guanylhydrazones of (Hetero)Aryl Methyl Ketones: Structure and Reaction with Acetic Anhydride. Monatshefte für Chemie, 1999, 130, 899-913.	0.9	7
149	Synthesis and odor of chiral partial structures of β -vetivone. I. Chirality, 1999, 11, 14-20.	1.3	7
150	Unambiguous Assignment of the 1H- and 13C-NMR Spectra of Propafenone and a Thiophene Analogue. Molecules, 2001, 6, 796-802.	1.7	7
151	A one-step synthesis of pyrazolone. MolBank, 2006, 2006, M464.	0.2	7
152	Synthesis of 1,3,4-tetrahydrospiro[chromene-2,2'-indoles] as a new class of ultrafast light-driven molecular switch. Tetrahedron, 2013, 69, 9309-9315.	1.0	7
153	An unusual thionyl chloride-promoted C-C bond formation to obtain 4,4'-bipyrazolones. Beilstein Journal of Organic Chemistry, 2018, 14, 1287-1292.	1.3	7
154	2-Amino-4-aryl-1-arylideneaminoimidazoles and Acylation Products: A Multinuclear (1 H, 13 C, 15 N) NMR Study. Monatshefte für Chemie, 2004, 135, 173-184.	0.9	6
155	Synthesis and antiproliferative activity of new cytotoxic azanaphthoquinone pyrrolo-annelated derivatives: Part II. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 3117-3121.	1.0	6
156	Synthesis and in Silico Evaluation of Novel Compounds for PET-Based Investigations of the Norepinephrine Transporter. Molecules, 2015, 20, 1712-1730.	1.7	6
157	Anti-inflammatory Effects of Compounds from Polygonum odoratum. Natural Product Communications, 2016, 11, 1934578X1601101.	0.2	6
158	Synthesis, Biological, and Computational Evaluation of Antagonistic, Chiral Hydrobenzoin Esters of Arecaidine Targeting mAChR M1. Pharmaceuticals, 2020, 13, 437.	1.7	6
159	Pyridazines, LVIII: 1-Phenyl-1-pyridazinyl-2-substituted ethenes, synthesis and configuration. Monatshefte für Chemie, 1991, 122, 1055-1061.	0.9	5
160	Synthesis and odor of chiral partial structures of β -vetivone, Part 2. Chirality, 1999, 11, 133-138.	1.3	5
161	Synthesis and antiproliferative activity of new cytotoxic tri- and tetraazabenzofluorene-5,6-dione derivatives. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 5264-5266.	1.0	5
162	H-Bond activated glycosylation of nucleobases: implications for prebiotic nucleoside synthesis. RSC Advances, 2014, 4, 3158-3161.	1.7	5

#	ARTICLE	IF	CITATIONS
163	Synthesis of tetrasubstituted pyrazoles containing pyridinyl substituents. <i>Beilstein Journal of Organic Chemistry</i> , 2017, 13, 895-902.	1.3	5
164	Consecutive C1-Homologation / Displacement Strategy for Converting Thiosulfonates into O,S-Homothioacetals. <i>Advanced Synthesis and Catalysis</i> , 2020, 362, 5444-5449.	2.1	5
165	N-Substituted 5,5-Dimethyl-2,5-dihydro-4H-isoindol-4-ones: Synthesis and NMR-Investigation. <i>Heterocycles</i> , 1996, 43, 1911.	0.4	5
166	Pyridazines, LXXII: On the Synthesis of Azinium and Diazinium Compounds Structurally Related to Pyridazomycin. <i>Archiv Der Pharmazie</i> , 1995, 328, 307-312.	2.1	4
167	Studies on ring opening reactions of β -lactams. <i>Tetrahedron</i> , 1997, 53, 8439-8446.	1.0	4
168	A Simple Synthesis of 6-Phenylpyrano[2,3-c]pyrazol-4(1H)-ones. <i>Synthesis</i> , 2005, 2005, 2583-2589.	1.2	4
169	Synthesis of 5-acyl-6-[2-hydroxy-3-(amino)propylamino]-1,3-dialkyl-1H-pyrimidine-2,4-diones. <i>Organic and Biomolecular Chemistry</i> , 2005, 3, 3958.	1.5	4
170	Synthesis of Mono- and Dibromo-Derivatives of 1-Phenylpyrazol-3-ol. <i>MolBank</i> , 2007, 2007, M551.	0.2	4
171	Heterocyclic analogs of xanthonines: 5,6-fused 3-methyl-1-phenylpyrano[2,3-c]pyrazol-4(1H)-thiones synthesis and NMR (¹ H, ¹³ C, ¹⁵ N) data. <i>Magnetic Resonance in Chemistry</i> , 2010, 48, 476-482.	1.1	4
172	Synthesis and reactions of 1-hydroxy-9,9a-dihydro-1H-imidazo[1,2-a]indol-2-(3H)-ones. <i>Tetrahedron</i> , 2011, 67, 3945-3953.	1.0	4
173	Azido derivatives of cellobiose: oxidation at C1 with cellobiose dehydrogenase from <i>Sclerotium rolfsii</i> . <i>Carbohydrate Research</i> , 2013, 382, 86-94.	1.1	4
174	Synthesis and biological evaluation of new cytotoxic indazolo[4,3-g]isoquinolinone derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 1846-1852.	1.0	4
175	The use of the Comins-Meyers Amide in Synthetic Chemistry: An Overview. <i>Natural Product Communications</i> , 2016, 11, 1934578X1601101.	0.2	4
176	2-Pyrazolin-5-ones Bearing a Basic Dialkylaminoalkyl Substituent at the N1-Position: Preparation and NMR Spectroscopic Investigations. <i>Heterocycles</i> , 2008, 75, 2035.	0.4	4
177	Synthesis of pyridyl substituted pyrazolo[4,3-c]pyridines as potential inhibitors of protein kinases. <i>Arkivoc</i> , 2014, 2014, 135-149.	0.3	4
178	Straightforward synthesis of bench-stable heteroatom-centered difluoromethylated entities via controlled nucleophilic transfer from activated TMSCHF ₂ . <i>Chemical Communications</i> , 2022, 58, 5761-5764.	2.2	4
179	Design, Synthesis, and Biological Evaluation of 4,4-TM-Difluorobenzhydryl Carbamates as Selective M1 Antagonists. <i>Pharmaceuticals</i> , 2022, 15, 248.	1.7	4
180	Synthesis, Biological Evaluation, and Docking Studies of Antagonistic Hydroxylated Arecaidine Esters Targeting mAChRs. <i>Molecules</i> , 2022, 27, 3173.	1.7	4

#	ARTICLE	IF	CITATIONS
181	A Method for the Synthesis of 2,3-Disubstituted 2,3-Dihydrobenzofurans. Monatshefte für Chemie, 2000, 131, 0375-0382.	0.9	3
182	On the Synthesis and Reactivity of 4-(Oxiran-2-ylmethoxy)cinnoline: Targeting a Cinnoline Analogue of Propranolol. Scientia Pharmaceutica, 2008, 76, 19-32.	0.7	3
183	3-Methyl-1-phenyl-1H-pyrazol-5-yl 2-Bromo-3-furan-carboxylate. MolBank, 2009, 2009, M603.	0.2	3
184	4-Bromo-3-methoxy-1-phenyl-1H-pyrazole. MolBank, 2009, 2009, M639.	0.2	3
185	(2E)-3-(3-Methoxy-1-phenyl-1H-pyrazol-4-yl)-2-propenal. MolBank, 2009, 2009, M644.	0.2	3
186	Reactions and Tautomeric Behavior of 1-(2-Pyridinyl)-1H-pyrazol-5-ols. Heterocycles, 2011, 83, 1567.	0.4	3
187	Halogen-impacted Reactivity in Lithium Carbenoid Mediated Homologations of Imine Surrogates: Direct Assembly of bis-trifluoromethyl-diketiminates and the Dual Role of LiCH ₂ I. Angewandte Chemie, 2020, 132, 21038-21043.	1.6	3
188	Straightforward and direct access to α -seleno- amines and sulfonylamides via the controlled addition of phenylselenomethyl lithium (LiCH ₂ SePh) to imines. Tetrahedron, 2020, 76, 131220.	1.0	3
189	Direct and straightforward transfer of C1 functionalized synthons to phosphorous electrophiles for accessing gem-P-containing methanes. Organic and Biomolecular Chemistry, 2021, 19, 2425-2429.	1.5	3
190	4,4'-bis-((2-Chlorophenyl)methylene)bis[1-phenyl-3-(trifluoromethyl)-1H-pyrazol-5-ol]. MolBank, 2009, 2009, M605.	0.2	2
191	5-Chloro-4-iodo-1,3-dimethyl-1H-pyrazole. MolBank, 2009, 2009, M620.	0.2	2
192	(2-Chlorophenyl)-3-methylchromeno[2,3-c]pyrazol-4(1H)-one. MolBank, 2010, 2010, M661.	0.2	2
193	4-[[[1-Phenyl-1H-pyrazol-3-yl]oxy]methyl]-1,3-dioxolan-2-one. MolBank, 2012, 2012, M786.	0.2	2
194	Pd-assisted Cross-coupling Reactions with 4-Chlorocinnoline. Journal of Heterocyclic Chemistry, 2013, 50, 141-144.	1.4	2
195	One-pot synthesis of polycyclic heterocyclic compounds by condensation of 1-carbamoylmethyl-2,3,3-trimethyl-3H-indolium salts with pyridine-2, 3, and 4- and quinoline-4-carboxaldehydes. Tetrahedron, 2018, 74, 3679-3690.	1.0	2
196	Synthesis of 10-methyl-8,10-diazabicyclo[4.3.1]decane as a new building block for nicotinic modulators. Arkivoc, 2013, 2013, 240-250.	0.3	2
197	Preparation of Amino- and Carboxy-Protected L- α -Amino- β -iodocarboxylic Acids. Archiv Der Pharmazie, 1995, 328, 367-370.	2.1	1
198	A ring-fission/C-C bond cleavage reaction with an N-alkyl-N-methyl-N-[(5-phenyl-1,2,4-oxadiazol-3-yl)methyl]amine. Tetrahedron, 2002, 58, 10417-10422.	1.0	1

#	ARTICLE	IF	CITATIONS
199	N-(3,4-Dichlorobenzyl)azoles—Investigations Regarding Synthesis, NMR-Spectroscopy and Affinity Towards Sisma-1 and Sigma-2 Receptors. <i>Scientia Pharmaceutica</i> , 2004, 72, 197-211.	0.7	1
200	Synthesis of 4,4'-(Cyclohexane-1,1-diyl)bis(1-methyl-1H-pyrazol-5-ol). <i>MolBank</i> , 2008, 2008, M569.	0.2	1
201	Synthesis and Ring Opening of Alkaloid-Type Compounds with a Novel Indolo[2,3-c][2]benzazepine Skeleton. <i>Synlett</i> , 2009, 2009, 3119-3122.	1.0	1
202	1-Phenylpyrazolo[4',3':5,6]pyrano[3,2-c]pyridine-4(1H)-thione. <i>MolBank</i> , 2010, 2010, M678.	0.2	1
203	Synthesis and NMR—Spectroscopic Investigations with 4—Chloroacetyl—phenylpyrazolin—ones. <i>Journal of Heterocyclic Chemistry</i> , 2018, 55, 132-137.	1.4	1
204	A ¹³ C chemical shifts study of iodopyrazoles: experimental results and relativistic and non-relativistic calculations. <i>Structural Chemistry</i> , 2021, 32, 925-937.	1.0	1
205	Unexpected scaffold rearrangement product of pirenzepine found in commercial samples. <i>Scientific Reports</i> , 2021, 11, 23397.	1.6	1
206	Synthesis of Substituted 3-Phenyl-6H-pyrazolo[4,3-d]isoxazoles (V) from Corresponding 4-Benzoyl-5-hydroxypyrazoles (I).. <i>ChemInform</i> , 2003, 34, no.	0.1	0
207	New 1-Substituted 4-Cinnamoyl-5-hydroxypyrazoles and Precursors Thereof: Synthesis, Ring Closure Reactions and NMR-Spectroscopic Investigations.. <i>ChemInform</i> , 2004, 35, no.	0.1	0
208	Spiro-Fused (C2)-Azirino-(C4)-pyrazolones, a New Heterocyclic System. Synthesis, Spectroscopic Studies and X-Ray Structure Analysis.. <i>ChemInform</i> , 2004, 35, no.	0.1	0
209	Synthesis and Detailed Spectroscopic Characterization of Two Novel N-(3-Acetyl-2-thienyl)acetamides. <i>MolBank</i> , 2006, 2006, M520.	0.2	0
210	5-Dimethylamino-1-phenylchromeno[2,3-c]pyrazol-4(1H)-one. <i>MolBank</i> , 2010, 2010, M706.	0.2	0
211	A One-Step Microwave-Assisted Synthetic Method for an O/S-Chemoselective Route to Derivatives of the First Adenosine A3 PET Radiotracer. <i>Molecules</i> , 2014, 19, 4076-4082.	1.7	0
212	2-Fluoro-N-methyl-N-((3S,4S)-4-[2-(trifluoromethyl)phenoxy]-3,4-dihydro-1H-isochromen-3-yl)methyl)ethanamine. <i>MolBank</i> , 2015, 2015, M858.	0.2	0
213	1-(3-Amino-1-phenylpropyl)-3-(2-fluorophenyl)-1,3-dihydro-2H-benzimidazol-2-one. <i>MolBank</i> , 2015, 2015, M867.	0.2	0
214	2-Fluoro-N-methyl-N-[(3S*,4S*)-4-(2-methylphenoxy)-3,4-dihydro-1H-isochromen-3-yl)methyl]ethanamine. <i>MolBank</i> , 2015, 2015, M862.	0.2	0
215	Consecutive and Selective Double Methylene Insertion of Lithium Carbenoids to Isothiocyanates: A Direct Assembly of Four—membered Sulfur—Containing Cycles. <i>Angewandte Chemie</i> , 0, , .	1.6	0
216	Synthesis and reactions of 1-amino-1,5,6,10b-tetrahydroimidazo[2,1-a]isoquinolin-2(3H)-ones. <i>Arkivoc</i> , 2009, 2009, 48-62.	0.3	0

#	ARTICLE	IF	CITATIONS
217	Imidazoquinazolinodiones " New Results. Heterocycles, 1999, 51, 1597.	0.4	0