

Franz Bracher

List of Publications by Year in descending order

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179
papers

4,572
citations

126907

33
h-index

138484

58
g-index

197
all docs

197
docs citations

197
times ranked

6936
citing authors

#	ARTICLE	IF	CITATIONS
1	Farm dust and endotoxin protect against allergy through A20 induction in lung epithelial cells. <i>Science</i> , 2015, 349, 1106-1110.	12.6	483
2	Specific CLK Inhibitors from a Novel Chemotype for Regulation of Alternative Splicing. <i>Chemistry and Biology</i> , 2011, 18, 67-76.	6.0	173
3	Accumulation of 8,9-unsaturated sterols drives oligodendrocyte formation and remyelination. <i>Nature</i> , 2018, 560, 372-376.	27.8	170
4	High susceptibility to fatty liver disease in two-pore channel 2-deficient mice. <i>Nature Communications</i> , 2014, 5, 4699.	12.8	164
5	Multiresidue analytical method using dispersive solid-phase extraction and gas chromatography/ion trap mass spectrometry to determine pharmaceuticals in whole blood. <i>Journal of Chromatography A</i> , 2006, 1135, 19-26.	3.7	144
6	A small molecule restores function to TRPML1 mutant isoforms responsible for mucopolidosis type IV. <i>Nature Communications</i> , 2014, 5, 4681.	12.8	125
7	Benzodiazepines and benzotriazepines as protein interaction inhibitors targeting bromodomains of the BET family. <i>Bioorganic and Medicinal Chemistry</i> , 2012, 20, 1878-1886.	3.0	112
8	Agonist-mediated switching of ion selectivity in TPC2 differentially promotes lysosomal function. <i>ELife</i> , 2020, 9, .	6.0	108
9	Sterol Biosynthesis and Azole Tolerance Is Governed by the Opposing Actions of SrbA and the CCAAT Binding Complex. <i>PLoS Pathogens</i> , 2016, 12, e1005775.	4.7	95
10	Functional Role and Therapeutic Potential of the Pim-1 Kinase in Colon Carcinoma. <i>Neoplasia</i> , 2013, 15, 783-IN28.	5.3	84
11	<i>Schistosoma mansoni</i> Sirtuins: Characterization and Potential as Chemotherapeutic Targets. <i>PLoS Neglected Tropical Diseases</i> , 2013, 7, e2428.	3.0	77
12	Synthesis and Biological Investigation of Phenothiazine-Based Benzhydroxamic Acids as Selective Histone Deacetylase 6 Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 1138-1166.	6.4	75
13	A human genome-wide loss-of-function screen identifies effective chikungunya antiviral drugs. <i>Nature Communications</i> , 2016, 7, 11320.	12.8	72
14	Selective agonist of TRPML2 reveals direct role in chemokine release from innate immune cells. <i>ELife</i> , 2018, 7, .	6.0	71
15	Novel 3-Arylideneindolin-2-ones as Inhibitors of NAD ⁺ -Dependent Histone Deacetylases (Sirtuins). <i>Journal of Medicinal Chemistry</i> , 2010, 53, 1383-1386.	6.4	69
16	7,8-Dichloro-1-oxo-1 ^H -carbolines as a Versatile Scaffold for the Development of Potent and Selective Kinase Inhibitors with Unusual Binding Modes. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 403-413.	6.4	64
17	Total Synthesis of the Pentacyclic Alkaloid Ascidiemin. <i>Heterocycles</i> , 1989, 29, 2093.	0.7	62
18	Azasteroids as antifungals. <i>Steroids</i> , 2003, 68, 587-594.	1.8	62

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19	Polycyclische aromatische Alkaloide, I. Synthese von Cleistopholin und Sampangin. Liebigs Annalen Der Chemie, 1989, 1989, 87-88.	0.8	56
20	1,2-Carbolin-Alkaloide, I. Synthese von 1-Aryl- und 1-Alkenyl-1,2-Carbolinen durch Palladium-katalysierte Kupplungsreaktionen. Liebigs Annalen Der Chemie, 1992, 1992, 1315-1319.	0.8	55
21	An Antifungal Benzimidazole Derivative Inhibits Ergosterol Biosynthesis and Reveals Novel Sterols. Antimicrobial Agents and Chemotherapy, 2015, 59, 6296-6307.	3.2	52
22	Selectivity Profiling and Biological Activity of Novel 1,2-Carbolines as Potent and Selective DYRK1 Kinase Inhibitors. PLoS ONE, 2015, 10, e0132453.	2.5	49
23	Development of Selective CBP/P300 Benzoxazepine Bromodomain Inhibitors. Journal of Medicinal Chemistry, 2016, 59, 8889-8912.	6.4	49
24	Antifungal drug testing by combining minimal inhibitory concentration testing with target identification by gas chromatography-mass spectrometry. Nature Protocols, 2017, 12, 947-963.	12.0	48
25	Saludimerines A and B, Novel-Type Dimeric Alkaloids with Stereogenic Centers and Configurationally Semistable Biaryl Axes. Journal of Organic Chemistry, 2004, 69, 8602-8608.	3.2	44
26	Fast and easy in vitro screening assay for cholesterol biosynthesis inhibitors in the post-squalene pathway. Steroids, 2007, 72, 633-642.	1.8	44
27	1,9-Dimetalated Δ^5 -carbolines. Versatile building blocks for the total synthesis of Alkaloids. Tetrahedron, 1994, 50, 12329-12336.	1.9	43
28	A convenient cellular assay for the identification of the molecular target of ergosterol biosynthesis inhibitors and quantification of their effects on total ergosterol biosynthesis. Steroids, 2013, 78, 483-493.	1.8	41
29	Inhibition of the SR Protein-Phosphorylating CLK Kinases of Plasmodium falciparum Impairs Blood Stage Replication and Malaria Transmission. PLoS ONE, 2014, 9, e105732.	2.5	39
30	Inhibition of Δ^5 -dehydrocholesterol reductase activates pro-resolving lipid mediator biosynthesis and inflammation resolution. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 20623-20634.	7.1	38
31	Antifungal defense of probiotic Lactobacillus rhamnosus GG is mediated by blocking adhesion and nutrient depletion. PLoS ONE, 2017, 12, e0184438.	2.5	38
32	Discovery of a Selective Allosteric Inhibitor Targeting Macrodomain 2 of Polyadenosine-Diphosphate-Ribose Polymerase 14. ACS Chemical Biology, 2017, 12, 2866-2874.	3.4	37
33	Gene editing and synthetically accessible inhibitors reveal role for TPC2 in HCC cell proliferation and tumor growth. Cell Chemical Biology, 2021, 28, 1119-1131.e27.	5.2	36
34	1,2-Carbolin-Alkaloide, II Tributyl(1-Ethoxyvinyl)stannan als C ₂ -Baustein für die Synthese von 1,2-Carbolin-Alkaloiden. Liebigs Annalen Der Chemie, 1993, 1993, 837-839.	0.8	33
35	Side chain azasteroids and thiaasteroids as sterol methyltransferase inhibitors in ergosterol biosynthesis. Bioorganic and Medicinal Chemistry, 2009, 17, 8123-8137.	3.0	32
36	Inhibition of endothelial Cdk5 reduces tumor growth by promoting non-productive angiogenesis. Oncotarget, 2016, 7, 6088-6104.	1.8	32

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37	Lathosterol side chain amides – A new class of human lathosterol oxidase inhibitors. <i>Steroids</i> , 2008, 73, 299-308.	1.8	29
38	Inhibition of NAD ⁺ -dependent histone deacetylases (sirtuins) causes growth arrest and activates both apoptosis and autophagy in the pathogenic protozoan <i>Trypanosoma cruzi</i> . <i>Parasitology</i> , 2014, 141, 814-825.	1.5	29
39	Total Synthesis of the Marine Pyridoacridine Alkaloid Demethyldeoxyamphimedine. <i>Journal of Organic Chemistry</i> , 2014, 79, 7239-7242.	3.2	29
40	Polycyclische aromatische Alkaloide, 2.Mitt. Synthese von Onychin und Eupolauridin. <i>Archiv Der Pharmazie</i> , 1989, 322, 293-294.	4.1	28
41	Total Synthesis of the Indolizinium Alkaloid Ficuseptine. <i>European Journal of Organic Chemistry</i> , 2002, 2002, 2288.	2.4	27
42	A New Class of Selective and Potent 7-Dehydrocholesterol Reductase Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 7614-7622.	6.4	27
43	A gas chromatography–mass spectrometry-based whole-cell screening assay for target identification in distal cholesterol biosynthesis. <i>Nature Protocols</i> , 2019, 14, 2546-2570.	12.0	27
44	DFG-1 Residue Controls Inhibitor Binding Mode and Affinity, Providing a Basis for Rational Design of Kinase Inhibitor Selectivity. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 10224-10234.	6.4	26
45	Discovery of a novel allosteric inhibitor scaffold for polyadenosine-diphosphate-ribose polymerase 14 (PARP14) macrodomain 2. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 2965-2972.	3.0	25
46	Resveratrol, lunularin and dihydroresveratrol do not act as caloric restriction mimetics when administered intraperitoneally in mice. <i>Scientific Reports</i> , 2019, 9, 4445.	3.3	25
47	Lung emphysema and impaired macrophage elastase clearance in mucolipin 3 deficient mice. <i>Nature Communications</i> , 2022, 13, 318.	12.8	25
48	KH-TFMDI, a novel sirtuin inhibitor, alters the cytoskeleton and mitochondrial metabolism promoting cell death in <i>Leishmania amazonensis</i> . <i>Apoptosis: an International Journal on Programmed Cell Death</i> , 2017, 22, 1169-1188.	4.9	24
49	Hetero Analogues of the Antimicrobial Alkaloids Cleistopholine and Sampangine. <i>Archiv Der Pharmazie</i> , 2007, 340, 429-433.	4.1	23
50	Estradiol analogs attenuate autophagy, cell migration and invasion by direct and selective inhibition of TRPML1, independent of estrogen receptors. <i>Scientific Reports</i> , 2021, 11, 8313.	3.3	23
51	Revised Structure of the Alkaloid Drymaritin. <i>Journal of Natural Products</i> , 2009, 72, 1908-1910.	3.0	22
52	A new approach to 1-substituted $\hat{2}$ -carbolines and isoquinolines utilizing tributyl[(Z)-2-ethoxyvinyl]stannane as a C-3,C-4 building block. <i>Tetrahedron</i> , 2016, 72, 837-845.	1.9	22
53	Regulation of influenza A virus mRNA splicing by CLK1. <i>Antiviral Research</i> , 2019, 168, 187-196.	4.1	21
54	The cytochrome <i>b₅</i> CyB is regulated by iron availability and is crucial for azole resistance in <i>A. fumigatus</i> . <i>Metallomics</i> , 2017, 9, 1655-1665.	2.4	20

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55	A convenient approach to the canthinone ring system: Total synthesis of the alkaloids tuboflavine and norisotuboflavine. <i>Journal of Heterocyclic Chemistry</i> , 2009, 46, 770-773.	2.6	19
56	Cytotoxic ring A-modified steroid analogues derived from Grundmann's ketone. <i>European Journal of Medicinal Chemistry</i> , 2011, 46, 3227-3236.	5.5	19
57	A divergent approach to benzylisoquinoline-type and oxoaporphine alkaloids via regioselective direct ring metalation of alkoxy isoquinolines. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 7664-7672.	2.8	19
58	Polycyclische Aromatische Alkaloide, V. Synthese von 2-Bromleptoclinidinon. <i>Liebigs Annalen Der Chemie</i> , 1990, 1990, 205-206.	0.8	18
59	Polycyclic Aromatic Alkaloids, 8. The Structure of Neocalliactine Acetate – Proof by Total Synthesis. <i>Liebigs Annalen Der Chemie</i> , 1992, 1992, 1205-1207.	0.8	18
60	Synthesis of Desaza Analogues of Annomontine and Canthinone Alkaloids. <i>Archiv Der Pharmazie</i> , 2015, 348, 125-131.	4.1	18
61	Synthesis and Biological Evaluation of Novel <i>N</i> -Alkyl Tetra- and Decahydroisoquinolines: Novel Antifungals that Target Ergosterol Biosynthesis. <i>Archiv Der Pharmazie</i> , 2014, 347, 283-290.	4.1	17
62	Synthesis of the Azaoxoaporphine Alkaloid Sampangine and Ascididemin-type Pyridoacridines through TMPMgCl·LiCl-Mediated Ring Closure. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 1302-1308.	2.4	17
63	New Perspectives in the Chemistry of Marine Pyridoacridine Alkaloids. <i>Marine Drugs</i> , 2016, 14, 26.	4.6	17
64	Pharmacokinetic Enhancers (Boosters) – Escort for Drugs against Degrading Enzymes and Beyond. <i>Scientia Pharmaceutica</i> , 2018, 86, 43.	2.0	17
65	A modular approach to the bisbenzylisoquinoline alkaloids tetrandrine and isotetrandrine. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 3047-3068.	2.8	17
66	$\hat{1}^2$ -Carboline Alkaloids, V: Total Synthesis of the Antimicrobial Marine Alkaloid Eudistomin T. $\hat{1}^2$ -Carboline-Alkaloide, 5. Mitt.: Totalsynthese des marinen Alkaloides Eudistomin T. <i>Archiv Der Pharmazie</i> , 1994, 327, 121-122.	4.1	16
67	Synthesis of 3-alkylpyridines. Part 2. Synthesis of both enantiomers of niphatesine C2. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1995, , 2323.	0.9	16
68	Aminopropylindenes derived from Grundmann's ketone as a novel chemotype of oxidosqualene cyclase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2013, 63, 758-764.	5.5	16
69	From Lead to Drug Utilizing a Mannich Reaction: The Topotecan Story. <i>Archiv Der Pharmazie</i> , 2017, 350, e1600236.	4.1	16
70	Phytotherapy Adds to the Therapeutic Armamentarium for the Treatment of Mild-To-Moderate Lower Urinary Tract Symptoms in Men. <i>Urologia Internationalis</i> , 2020, 104, 333-342.	1.3	16
71	Short Total Synthesis of the Marine Alkaloid Subarine. <i>Scientia Pharmaceutica</i> , 2009, 77, 1-7.	2.0	15
72	N-Methylation of Aromatic Amines and N-Heterocycles under Acidic Conditions with the TTT (1,3,5-Trioxane – Triethylsilane – Trifluoroacetic Acid) System. <i>Synthesis</i> , 2015, 47, 3333-3338.	2.3	15

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73	Farm dust reduces viral load in human bronchial epithelial cells by increasing barrier function and antiviral responses. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 141, 1949-1952.e8.	2.9	15
74	Sterol Composition of Clinically Relevant Mucorales and Changes Resulting from Posaconazole Treatment. <i>Molecules</i> , 2018, 23, 1218.	3.8	15
75	The Putative Caloric Restriction Mimetic Resveratrol has Moderate Impact on Insulin Sensitivity, Body Composition, and the Metabolome in Mice. <i>Molecular Nutrition and Food Research</i> , 2020, 64, e1901116.	3.3	15
76	First total synthesis of the 2,7-naphthyridine alkaloid neozeylanine and unexpected formation of isoquinolines from ethoxyvinyl pyridines. <i>Liebigs Annalen</i> , 1995, 1995, 645-647.	0.8	14
77	Alkaloids from <i>Croton flavens</i> and their affinities to GABA-receptors. <i>Natural Product Research</i> , 2003, 17, 437-440.	1.8	14
78	Total Syntheses of the Chlorinated Carboline Alkaloids Bauerine A, B, and C. <i>Synthetic Communications</i> , 2007, 37, 1273-1280.	2.1	14
79	Analysis and Experimental Inhibition of Distal Cholesterol Biosynthesis. <i>Chromatographia</i> , 2015, 78, 343-358.	1.3	14
80	New chemotype of selective and potent inhibitors of human delta 24-dehydrocholesterol reductase. <i>European Journal of Medicinal Chemistry</i> , 2017, 140, 305-320.	5.5	14
81	Comparison of Strategies for the Determination of Sterol Sulfates via GC-MS Leading to a Novel Deconjugation-Derivatization Protocol. <i>Molecules</i> , 2019, 24, 2353.	3.8	14
82	Identification of the subtype-selective Sirt5 inhibitor balsalazide through systematic SAR analysis and rationalization via theoretical investigations. <i>European Journal of Medicinal Chemistry</i> , 2020, 206, 112676.	5.5	14
83	Polycyclische aromatische Alkaloide, 3. Mitt.: Synthese von Perlolidin Polycyclic Aromatic Alkaloids, III: Synthesis of Perlolidine. <i>Archiv Der Pharmazie</i> , 1989, 322, 511-512.	4.1	13
84	First synthesis of the benzo[<i>a</i>]pyrido[2,3- <i>b</i>]pyrrolo[2,1- <i>a</i>][2,7]naphthyridine ring system. <i>Journal of Heterocyclic Chemistry</i> , 1993, 30, 157-159.	2.6	13
85	Short and Efficient Approach Towards Macrocyclic Lactones Based on a Sonogashira Reaction. <i>Archiv Der Pharmazie</i> , 2005, 338, 605-608.	4.1	13
86	First Total Synthesis of the 2,7-Naphthyridine Alkaloids Lophocladine A and B. <i>Archiv Der Pharmazie</i> , 2006, 339, 677-679.	4.1	13
87	Regioselective homolytic substitution of benzo[<i>c</i>][2,7]naphthyridines. <i>Tetrahedron</i> , 2012, 68, 4693-4700.	1.9	13
88	A novel approach to ring A analogues of the marine pyridoacridine alkaloid ascididemin. <i>Tetrahedron</i> , 2013, 69, 9857-9864.	1.9	13
89	Discovery of lipophilic two-pore channel agonists. <i>FEBS Journal</i> , 2020, 287, 5284-5293.	4.7	13
90	Chemical and pharmacological characterization of the TRPML calcium channel blockers ML-S11 and ML-S13. <i>European Journal of Medicinal Chemistry</i> , 2021, 210, 112966.	5.5	13

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91	Dehydrocholesterol Reductase 24 (DHCR24): Medicinal Chemistry, Pharmacology and Novel Therapeutic Options. <i>Current Medicinal Chemistry</i> , 2022, 29, 4005-4025.	2.4	13
92	Polycyclische aromatische Alkaloide, 7. Mitt.: Studien zur Struktur von Dielsin. <i>Archiv Der Pharmazie</i> , 1992, 325, 645-648.	4.1	12
93	Î²-Carbolin-Alkaloide, IV. â€“ Synthese von 1-Alkyl-Î²-carbolinen und Strukturrevision von Lycii-Alkaloid I. <i>Liebigs Annalen Der Chemie</i> , 1993, 1993, 1335-1337.	0.8	12
94	Î²-Carboline alkaloids 9 [1]. Total synthesis of the Î²-carboline alkaloids arenarine A and (Â±)arenarine B. <i>Journal of Heterocyclic Chemistry</i> , 2004, 41, 173-176.	2.6	12
95	Stereoselective synthesis of a new class of potent and selective inhibitors of human Î² ^{8,7} -sterol isomerase. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 1925-1943.	3.0	12
96	New approaches to the synthesis of canthin-4-one alkaloids and synthetic analogues. <i>Tetrahedron</i> , 2015, 71, 4640-4646.	1.9	12
97	Synthesis and Structure-Activity Relationships of Novel Benzylamine-Type Antifungals as Butenafine-Related Antimycotics. <i>Archiv Der Pharmazie</i> , 2017, 350, 1600342.	4.1	12
98	Isoquinoline-based biaryls as a robust scaffold for microtubule inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2020, 186, 111865.	5.5	12
99	Fungal sterol C22-desaturase is not an antimycotic target as shown by selective inhibitors and testing on clinical isolates. <i>Steroids</i> , 2015, 101, 1-6.	1.8	11
100	Alterations on growth and cell organization of <i>Giardia intestinalis</i> trophozoites after treatment with KH-TFMDI, a novel class III histone deacetylase inhibitor. <i>International Journal of Medical Microbiology</i> , 2019, 309, 130-142.	3.6	11
101	Miniaturized multiresidue method for the analysis of pesticides and persistent organic pollutants in non-target wildlife animal liver tissues using GC-MS/MS. <i>Chemosphere</i> , 2021, 279, 130434.	8.2	11
102	Screening Health-Promoting Compounds for Their Capacity to Induce the Activity of FOXO3. <i>Journals of Gerontology - Series A Biological Sciences and Medical Sciences</i> , 2022, 77, 1485-1493.	3.6	11
103	Lysosomal TRPML1 regulates mitochondrial function in hepatocellular carcinoma cells. <i>Journal of Cell Science</i> , 2022, 135, .	2.0	11
104	Unexpected ipso-Substitutions at the Î²-Carboline Nucleus. <i>Synthetic Communications</i> , 2003, 33, 3843-3850.	2.1	10
105	A Novel Approach to the Pyridoacridine Ring System: Synthesis of the Topoisomerase Inhibitor 13-Deazaascididemin. <i>Archiv Der Pharmazie</i> , 2012, 345, 822-826.	4.1	10
106	A new approach to monoprotected 1,4-benzodiazepines via a one-pot N-deprotection/reductive cyclization procedure. <i>Tetrahedron</i> , 2016, 72, 1668-1674.	1.9	10
107	Arylpiperidines as a new class of oxidosqualene cyclase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2016, 109, 13-22.	5.5	10
108	The yeast pantothenate kinase Cab1 is a master regulator of sterol metabolism and of susceptibility to ergosterol biosynthesis inhibitors. <i>Journal of Biological Chemistry</i> , 2019, 294, 14757-14767.	3.4	10

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109	How to Separate Kinase Inhibition from Undesired Monoamine Oxidase A Inhibition – The Development of the DYRK1A Inhibitor AnnH75 from the Alkaloid Harmine. <i>Molecules</i> , 2020, 25, 5962.	3.8	10
110	The Total Synthesis of Both Enantiomers of the Macrocyclic Lactone Zearalane. <i>European Journal of Organic Chemistry</i> , 2001, 2001, 4701.	2.4	9
111	New Substituted Isocoumarins and Dihydroisocoumarins and their Cytotoxic Activities. <i>Scientia Pharmaceutica</i> , 2011, 79, 21-30.	2.0	9
112	One-Pot Conversion of 1-Bromo- β -carboline and 1-Bromocarbazole into Pentacyclic Compounds by Suzuki Cross-Coupling Followed by Spontaneous Cyclization. <i>Synthesis</i> , 2014, 46, 893-898.	2.3	9
113	A divergent approach to the total synthesis of the marine pyridoacridine alkaloid eilatin and its synthetic isomer isoelatin. <i>Tetrahedron Letters</i> , 2015, 56, 1445-1447.	1.4	9
114	Cytotoxic Hybrids Between the Aromatic Alkaloids Bauerine C and Rutaecarpine. <i>Zeitschrift Fur Naturforschung - Section B Journal of Chemical Sciences</i> , 2007, 62, 1313-1316.	0.7	8
115	7-Aza-des-A-steroids with Antimicrobial and Cytotoxic Activity. <i>Scientia Pharmaceutica</i> , 2013, 81, 329-338.	2.0	8
116	Synthesis and Biological Evaluation of Novel Alkyl-Imidazolyl Carbinols and their Esters: Potent Antimycotics. <i>Scientia Pharmaceutica</i> , 2013, 81, 641-650.	2.0	8
117	A novel approach to oxoisoaporphine alkaloids via regioselective metalation of alkoxy isoquinolines. <i>Beilstein Journal of Organic Chemistry</i> , 2017, 13, 1564-1571.	2.2	8
118	Synthesis, biological evaluation and toxicity of novel tetrandrine analogues. <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112810.	5.5	8
119	New polycyclic ring systems derived from canthinone. <i>Journal of Heterocyclic Chemistry</i> , 2010, 47, 449-453.	2.6	7
120	The Gramine Route to Pyrido[4,3-b]indol-3-ones – Identification of a New Cytotoxic Lead. <i>Scientia Pharmaceutica</i> , 2011, 79, 59-68.	2.0	7
121	Synthesis and Antifungal Evaluation of Novel N-Alkyl Tetra- and Perhydroquinoline Derivatives. <i>Scientia Pharmaceutica</i> , 2015, 83, 1-14.	2.0	7
122	Traceless bond construction via rearrangement of N-Boc-N-allylhydrazones giving 1,1-disubstituted olefins. <i>Tetrahedron</i> , 2015, 71, 2530-2539.	1.9	7
123	A short and divergent route to 2-alkenyl-4-quinolones. <i>Tetrahedron Letters</i> , 2018, 59, 3632-3635.	1.4	7
124	Development of a convenient method for the determination of dimethyl sulfoxide in lyophilised pharmaceuticals by static headspace gas chromatography-mass spectrometry. <i>Analytical Methods</i> , 2019, 11, 2119-2122.	2.7	7
125	Inhibition of Phytosterol Biosynthesis by Azasterols. <i>Molecules</i> , 2020, 25, 1111.	3.8	7
126	7,9,12b-Triazabenz[a]aceanthrylen-8-one, the First Representative of a Novel Pentacyclic Ring System and its Biological Activities. <i>Letters in Organic Chemistry</i> , 2013, 10, 568-572.	0.5	7

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127	Crotoflavol, A New Phenanthrene from <i>Croton flavens</i> . <i>Natural Product Research</i> , 2001, 15, 147-150.	0.4	6
128	1-Substituted β -Carboline-3-Carboxylates with high affinities to the Benzodiazepine recognition site. <i>Natural Product Research</i> , 2004, 18, 391-396.	1.8	6
129	Steroidomimetic Aminomethyl Spiroacetals as Novel Inhibitors of the Enzyme β -Sterol Isomerase in Cholesterol Biosynthesis. <i>Archiv Der Pharmazie</i> , 2014, 347, 108-122.	4.1	6
130	Canthinones as Novel Antibacterial Agents. <i>Archiv Der Pharmazie</i> , 2016, 349, 710-723.	4.1	6
131	In vitro production of reactive oxygen species (ROS) by sampangine. <i>Medicinal Chemistry Research</i> , 2017, 26, 1170-1175.	2.4	6
132	Chiral-pool synthesis of 1,2,4-trisubstituted 1,4-diazepanes as novel β 1 receptor ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 4778-4799.	3.0	6
133	Functionalization of 4-bromobenzo[<i>c</i>][2,7]naphthyridine via regioselective direct ring metalation. A novel approach to analogues of pyridoacridine alkaloids. <i>Beilstein Journal of Organic Chemistry</i> , 2019, 15, 2304-2310.	2.2	6
134	Expression, purification and crystallization of CLK1 kinase – A potential target for antiviral therapy. <i>Protein Expression and Purification</i> , 2020, 176, 105742.	1.3	6
135	Development of a human biomonitoring method for assessing the exposure to ethoxyquin in the general population. <i>Archives of Toxicology</i> , 2020, 94, 4209-4217.	4.2	6
136	Determination of multi pesticide residues in leaf and needle samples using a modified QuEChERS approach and gas chromatography-tandem mass spectrometry. <i>Analytical Methods</i> , 2021, 13, 1138-1146.	2.7	6
137	A new approach towards Ikimine a analogues. <i>Natural Product Research</i> , 2004, 18, 397-401.	1.8	5
138	A Convenient Conversion of Substituted Cyclohexenones into Aryl Methyl Ketones. <i>Synthesis</i> , 2012, 44, 2441-2447.	2.3	5
139	First total synthesis of the marine steroid alkaloid plakinamine B. <i>Tetrahedron</i> , 2014, 70, 1084-1090.	1.9	5
140	Reductive N-Arylethylolation of Aromatic Amines and N-Heterocycles with Enol Ethers. <i>Synthesis</i> , 2018, 50, 1323-1330.	2.3	5
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