

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	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
2	Effective chiral pool synthesis of both enantiomers of the TRPML inhibitor <i>trans</i> -ML-SI3. <i>Archiv Der Pharmazie</i> , 2022, 355, e2100362.	4.1	5
3	Dehydrocholesterol Reductase 24 (DHCR24): Medicinal Chemistry, Pharmacology and Novel Therapeutic Options. <i>Current Medicinal Chemistry</i> , 2022, 29, 4005-4025.	2.4	13
4	Lung emphysema and impaired macrophage elastase clearance in mucolipin 3 deficient mice. <i>Nature Communications</i> , 2022, 13, 318.	12.8	25
5	Using the yeast three-hybrid system for the identification of small molecule-protein interactions with the example of ethinylestradiol. <i>Biological Chemistry</i> , 2022, 403, 421-431.	2.5	1
6	Lysosomal TRPML1 regulates mitochondrial function in hepatocellular carcinoma cells. <i>Journal of Cell Science</i> , 2022, 135, .	2.0	11
7	In vitro effects of the 4-[(10H-phenothiazin-10-yl)methyl]-N-hydroxybenzamide on <i>Giardia intestinalis</i> trophozoites. <i>Acta Tropica</i> , 2022, 232, 106484.	2.0	4
8	Analysis of pesticide and persistent organic pollutant residues in German bats. <i>Chemosphere</i> , 2022, 305, 135342.	8.2	3
9	Development of hetero-triaryls as a new chemotype for subtype-selective and potent Sirt5 inhibition. <i>European Journal of Medicinal Chemistry</i> , 2022, 240, 114594.	5.5	3
10	Chemical and pharmacological characterization of the TRPML calcium channel blockers ML-SI1 and ML-SI3. <i>European Journal of Medicinal Chemistry</i> , 2021, 210, 112966.	5.5	13
11	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
12	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
13	DMSO as new, counterintuitive excipient for freeze-drying human keratinocytes. <i>European Journal of Pharmaceutical Sciences</i> , 2021, 160, 105746.	4.0	5
14	C81-evoked inhibition of the TNFR1-NF κ B pathway during inflammatory processes for stabilization of the impaired vascular endothelial barrier for leukocytes. <i>FASEB Journal</i> , 2021, 35, e21656.	0.5	3
15	Effective sample preparation procedure for the analysis of free neutral steroids, free steroid acids and sterol sulfates in different tissues by GC-MS. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2021, 211, 105880.	2.5	1
16	Gene editing and synthetically accessible inhibitors reveal role for TPC2 in HCC cell proliferation and tumor growth. <i>Cell Chemical Biology</i> , 2021, 28, 1119-1131.e27.	5.2	36
17	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
18	A Short Approach to N-Aryl-1,2,3,4-tetrahydroisoquinolines from N-(2-Bromobenzyl)anilines via a Reductive Amination/Palladium-Catalyzed Ethoxyvinylolation/Reductive N-Alkylation Sequence. <i>Synthesis</i> , 2021, 53, 1943-1954.	2.3	4

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19	Synthesis of highly substituted fluorenones via metal-free TBHP-promoted oxidative cyclization of 2-(aminomethyl)biphenyls. Application to the total synthesis of nobilone. <i>Beilstein Journal of Organic Chemistry</i> , 2021, 17, 2668-2679.	2.2	5
20	The ethoxycarbonyl group as both activating and protective group in N-acyl-Pictetâ€“Spengler reactions using methoxystyrenes. A short approach to racemic 1-benzyltetrahydroisoquinoline alkaloids. <i>Beilstein Journal of Organic Chemistry</i> , 2021, 17, 2716-2725.	2.2	2
21	Synthesis, Biological Evaluation, and Structureâ€“Activity Relationships of 4-Aminopiperidines as Novel Antifungal Agents Targeting Ergosterol Biosynthesis. <i>Molecules</i> , 2021, 26, 7208.	3.8	4
22	A short divergent approach to highly substituted carbazoles and Î²-carbolines via in situ-generated diketoindoles. <i>Tetrahedron Letters</i> , 2020, 61, 151597.	1.4	2
23	Greener aromatic antioxidants for aviation and beyond. <i>Sustainable Energy and Fuels</i> , 2020, 4, 2153-2163.	4.9	4
24	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
25	Isoquinoline-based biaryls as a robust scaffold for microtubule inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2020, 186, 111865.	5.5	12
26	Pharmacological characterization of high-affinity Î²1 receptor ligands with spirocyclic thienopyran and thienofuran scaffold. <i>Journal of Pharmacy and Pharmacology</i> , 2020, 72, 236-248.	2.4	3
27	Synthesis of Seco â€“Analogues of the DHCR24 Inhibitor SHâ€“42. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 6270-6288.	2.4	1
28	Synthesis, biological evaluation and toxicity of novel tetrandrine analogues. <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112810.	5.5	8
29	Expression, purification and crystallization of CLK1 kinase â€“ A potential target for antiviral therapy. <i>Protein Expression and Purification</i> , 2020, 176, 105742.	1.3	6
30	Racemic total synthesis and evaluation of the biological activities of the isoquinolineâ€“benzylisoquinoline alkaloid muraricine. <i>Archiv Der Pharmazie</i> , 2020, 353, 2000106.	4.1	5
31	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
32	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
33	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
34	How to Separate Kinase Inhibition from Undesired Monoamine Oxidase A Inhibitionâ€“The Development of the DYRK1A Inhibitor AnnH75 from the Alkaloid Harmine. <i>Molecules</i> , 2020, 25, 5962.	3.8	10
35	Discovery of lipophilic twoâ€“pore channel agonists. <i>FEBS Journal</i> , 2020, 287, 5284-5293.	4.7	13
36	Inhibition of Phytosterol Biosynthesis by Azasterols. <i>Molecules</i> , 2020, 25, 1111.	3.8	7

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37	Human metabolism and urinary excretion kinetics of the UV filter Uvinul A plus [®] after a single oral or dermal dosage. <i>International Journal of Hygiene and Environmental Health</i> , 2020, 227, 113509.	4.3	3
38	A Novel Approach to Highly Substituted ¹² C- ¹³ C Carbolines via Reductive Ring Transformation of 2-acyl-3-isoxazolyloindoles. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 2708-2719.	2.4	5
39	Dimethylformamide Acetals and Brederick's Reagent as Building Blocks in Natural Products Total Synthesis. <i>Mini-Reviews in Organic Chemistry</i> , 2020, 17, 47-66.	1.3	5
40	A modular approach to the bisbenzylisoquinoline alkaloids tetrandrine and isotetrandrine. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 3047-3068.	2.8	17
41	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
42	Traceless Isoprenylation of Aldehydes via <i>N</i> -Boc- <i>N</i> -(1,1-dimethylallyl)hydrazones. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 3680-3687.	2.4	3
43	A versatile approach to 1-oxo-, 1-oxo-3,4-dihydro- and 1,3,4-trioxo isoquinoline alkaloids and first total synthesis of the dimeric 1-oxoisoquinoline alkaloids berbanine and berbidine. <i>Tetrahedron</i> , 2020, 76, 131150.	1.9	5
44	Agonist-mediated switching of ion selectivity in TPC2 differentially promotes lysosomal function. <i>ELife</i> , 2020, 9, .	6.0	108
45	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
46	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
47	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
48	Inhibition of ¹⁴ C-dehydrocholesterol reductase activates pro-resolving lipid mediator biosynthesis and inflammation resolution. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 20623-20634.	7.1	38
49	Novel access to 2-substituted quinolin-4-ones by nickel boride-mediated reductive ring transformation of 5-(2-nitrophenyl)isoxazoles. <i>Tetrahedron Letters</i> , 2019, 60, 151327.	1.4	4
50	Characterization of two new degradation products of atorvastatin calcium formed upon treatment with strong acids. <i>Beilstein Journal of Organic Chemistry</i> , 2019, 15, 2085-2091.	2.2	3
51	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
52	Regulation of influenza A virus mRNA splicing by CLK1. <i>Antiviral Research</i> , 2019, 168, 187-196.	4.1	21
53	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
54	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

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55	A validated UPLC-MS/MS method for the determination of urinary metabolites of Uvinul® A plus. <i>Analytical and Bioanalytical Chemistry</i> , 2019, 411, 8143-8152.	3.7	4
56	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
57	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
58	General Method for the Preparation of Indole-2-Weinreb Amides. <i>Synthesis</i> , 2019, 51, 757-768.	2.3	3
59	Blocking Lysosomal Two-Pore Channel 2 Function Inhibits Proliferation of Multidrug Resistant Leukemia Cells and Sensitizes Them to Vincristine Treatment. <i>Blood</i> , 2019, 134, 2081-2081.	1.4	3
60	A Short Synthesis of the Plant Alkaloid 4-Methyl-2,6-naphthyridine. <i>Letters in Organic Chemistry</i> , 2019, 16, 931-934.	0.5	5
61	Methods for Arylethylation of Amines and Heteroarenes. <i>SynOpen</i> , 2018, 02, 0096-0104.	1.7	2
62	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
63	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
64	Reductive N-Arylethylation of Aromatic Amines and N-Heterocycles with Enol Ethers. <i>Synthesis</i> , 2018, 50, 1323-1330.	2.3	5
65	Pharmacokinetic Enhancers (Boosters) – Escort for Drugs against Degrading Enzymes and Beyond. <i>Scientia Pharmaceutica</i> , 2018, 86, 43.	2.0	17
66	A short and divergent route to 2-alkenyl-4-quinolones. <i>Tetrahedron Letters</i> , 2018, 59, 3632-3635.	1.4	7
67	Aminomethylation/hydrogenolysis as an alternative to direct methylation of metalated isoquinolines – a novel total synthesis of the alkaloid 7-hydroxy-6-methoxy-1-methylisoquinoline. <i>Beilstein Journal of Organic Chemistry</i> , 2018, 14, 130-134.	2.2	5
68	Accumulation of 8,9-unsaturated sterols drives oligodendrocyte formation and remyelination. <i>Nature</i> , 2018, 560, 372-376.	27.8	170
69	Sterol Composition of Clinically Relevant Mucorales and Changes Resulting from Posaconazole Treatment. <i>Molecules</i> , 2018, 23, 1218.	3.8	15
70	Selective agonist of TRPML2 reveals direct role in chemokine release from innate immune cells. <i>ELife</i> , 2018, 7, .	6.0	71
71	Reversal of Chemoresistance in Leukemia Cells Using Synthetic Bisbenzylisoquinoline Derivatives. <i>Blood</i> , 2018, 132, 3504-3504.	1.4	0
72	In vitro production of reactive oxygen species (ROS) by sampangine. <i>Medicinal Chemistry Research</i> , 2017, 26, 1170-1175.	2.4	6

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73	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
74	Antifungal drug testing by combining minimal inhibitory concentration testing with target identification by gas chromatography-mass spectrometry. <i>Nature Protocols</i> , 2017, 12, 947-963.	12.0	48
75	Discovery of a Selective Allosteric Inhibitor Targeting Macrodomain 2 of Polyadenosine-Diphosphate-Ribose Polymerase 14. <i>ACS Chemical Biology</i> , 2017, 12, 2866-2874.	3.4	37
76	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
77	Chiral-pool synthesis of 1,2,4-trisubstituted 1,4-diazepanes as novel 5-HT _{1A} receptor ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 4778-4799.	3.0	6
78	Synthesis of highly substituted 3-arylideneindolin-2-ones. <i>Tetrahedron</i> , 2017, 73, 5668-5679.	1.9	3
79	The cytochrome <i>b₅</i> CybE is regulated by iron availability and is crucial for azole resistance in <i>A. fumigatus</i> . <i>Metalomics</i> , 2017, 9, 1655-1665.	2.4	20
80	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
81	From Lead to Drug Utilizing a Mannich Reaction: The Topotecan Story. <i>Archiv Der Pharmazie</i> , 2017, 350, e1600236.	4.1	16
82	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
83	Antifungal defense of probiotic <i>Lactobacillus rhamnosus</i> GG is mediated by blocking adhesion and nutrient depletion. <i>PLoS ONE</i> , 2017, 12, e0184438.	2.5	38
84	The Chemistry of the Canthin-4-ones. <i>Mini-Reviews in Organic Chemistry</i> , 2017, 14, 92-98.	1.3	3
85	Inhibition of endothelial Cdk5 reduces tumor growth by promoting non-productive angiogenesis. <i>Oncotarget</i> , 2016, 7, 6088-6104.	1.8	32
86	New Perspectives in the Chemistry of Marine Pyridoacridine Alkaloids. <i>Marine Drugs</i> , 2016, 14, 26.	4.6	17
87	Development of Selective CBP/P300 Benzoxazepine Bromodomain Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 8889-8912.	6.4	49
88	Canthin-4-ones as Novel Antibacterial Agents. <i>Archiv Der Pharmazie</i> , 2016, 349, 710-723.	4.1	6
89	A human genome-wide loss-of-function screen identifies effective chikungunya antiviral drugs. <i>Nature Communications</i> , 2016, 7, 11320.	12.8	72
90	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

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91	A new approach to 1-substituted Î²-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
92	Arylpiperidines as a new class of oxidosqualene cyclase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2016, 109, 13-22.	5.5	10
93	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
94	Triflimide-Catalysed Rearrangement of N-(Trimethylsilyl)allylhydrazones Results in the Formation of Vinylsilanes and Cyclopropanes. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 8024-8033.	2.4	4
95	Selectivity Profiling and Biological Activity of Novel Î²-Carbolines as Potent and Selective DYRK1 Kinase Inhibitors. <i>PLoS ONE</i> , 2015, 10, e0132453.	2.5	49
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 Antifungal Evaluation of Novel N-Alkyl Tetra- and Perhydroquinoline Derivatives. <i>Scientia Pharmaceutica</i> , 2015, 83, 1-14.	2.0	7
98	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
99	Synthesis of the Azaoxaporphine Alkaloid Sampangine and Ascidiemin-Type Pyridoacridines through TPMgCl-Li-Mediated Ring Closure. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 1302-1308.	2.4	17
100	A divergent approach to the total synthesis of the marine pyridoacridine alkaloid eilatin and its synthetic isomer isoeilatin. <i>Tetrahedron Letters</i> , 2015, 56, 1445-1447.	1.4	9
101	Synthesis of Desaza Analogues of Annomontine and Canthin-4-one Alkaloids. <i>Archiv Der Pharmazie</i> , 2015, 348, 125-131.	4.1	18
102	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
103	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
104	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
105	An Antifungal Benzimidazole Derivative Inhibits Ergosterol Biosynthesis and Reveals Novel Sterols. <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 6296-6307.	3.2	52
106	Farm dust and endotoxin protect against allergy through A20 induction in lung epithelial cells. <i>Science</i> , 2015, 349, 1106-1110.	12.6	483
107	Analysis and Experimental Inhibition of Distal Cholesterol Biosynthesis. <i>Chromatographia</i> , 2015, 78, 343-358.	1.3	14
108	A small molecule restores function to TRPML1 mutant isoforms responsible for mucopolipidosis type IV. <i>Nature Communications</i> , 2014, 5, 4681.	12.8	125

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109	One-Pot Conversion of 1-Bromo- $\hat{2}$ -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
110	Steroidomimetic Aminomethyl Spiroacetals as Novel Inhibitors of the Enzyme $\hat{8,7}$ -sterol Isomerase in Cholesterol Biosynthesis. <i>Archiv Der Pharmazie</i> , 2014, 347, 108-122.	4.1	6
111	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
112	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
113	First total synthesis of the marine steroid alkaloid plakinamine B. <i>Tetrahedron</i> , 2014, 70, 1084-1090.	1.9	5
114	Total Synthesis of the Marine Pyridoacridine Alkaloid Demethyldeoxyamphimedine. <i>Journal of Organic Chemistry</i> , 2014, 79, 7239-7242.	3.2	29
115	High susceptibility to fatty liver disease in two-pore channel 2-deficient mice. <i>Nature Communications</i> , 2014, 5, 4699.	12.8	164
116	Inhibition of the SR Protein-Phosphorylating CLK Kinases of <i>Plasmodium falciparum</i> Impairs Blood Stage Replication and Malaria Transmission. <i>PLoS ONE</i> , 2014, 9, e105732.	2.5	39
117	A novel approach to ring A analogues of the marine pyridoacridine alkaloid ascididemin. <i>Tetrahedron</i> , 2013, 69, 9857-9864.	1.9	13
118	Functional Role and Therapeutic Potential of the Pim-1 Kinase in Colon Carcinoma. <i>Neoplasia</i> , 2013, 15, 783-IN28.	5.3	84
119	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
120	Stereoselective synthesis of a new class of potent and selective inhibitors of human $\hat{8,7}$ -sterol isomerase. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 1925-1943.	3.0	12
121	A convenient cellular assay for the identification of the molecular target of ergosterol biosynthesis inhibitors and quantification of their effects on total ergosterol biosynthesis. <i>Steroids</i> , 2013, 78, 483-493.	1.8	41
122	<i>Schistosoma mansoni</i> Sirtuins: Characterization and Potential as Chemotherapeutic Targets. <i>PLoS Neglected Tropical Diseases</i> , 2013, 7, e2428.	3.0	77
123	7-Aza-des-A-steroids with Antimicrobial and Cytotoxic Activity. <i>Scientia Pharmaceutica</i> , 2013, 81, 329-338.	2.0	8
124	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
125	7,9,12b-Triazabenz[<i>a</i>]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
126	Isolation of the Pyrrolizidine Alkaloid Intermedine-N-oxide from <i>Cerithe glabra</i> and ab initio Calculation of its ¹³ C NMR Shifts. <i>Zeitschrift Fur Naturforschung - Section B Journal of Chemical Sciences</i> , 2012, 67, 411-416.	0.7	3

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127	A Convenient Conversion of Substituted Cyclohexenones into Aryl Methyl Ketones. <i>Synthesis</i> , 2012, 44, 2441-2447.	2.3	5
128	7,8-Dichloro-1-oxo- β -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
129	A New Class of Selective and Potent 7-Dehydrocholesterol Reductase Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 7614-7622.	6.4	27
130	A Novel Approach to the Pyridoacridine Ring System: Synthesis of the Topoisomerase Inhibitor 13 β -Deazaascididemin. <i>Archiv Der Pharmazie</i> , 2012, 345, 822-826.	4.1	10
131	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
132	Regioselective homolytic substitution of benzo[c][2,7]naphthyridines. <i>Tetrahedron</i> , 2012, 68, 4693-4700.	1.9	13
133	Specific CLK Inhibitors from a Novel Chemotype for Regulation of Alternative Splicing. <i>Chemistry and Biology</i> , 2011, 18, 67-76.	6.0	173
134	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
135	New Substituted Isocoumarins and Dihydroisocoumarins and their Cytotoxic Activities. <i>Scientia Pharmaceutica</i> , 2011, 79, 21-30.	2.0	9
136	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
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