

Mouad Alami

List of Publications by Year in descending order

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108
papers

4,167
citations

76294

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133188

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108
all docs

108
docs citations

108
times ranked

3441
citing authors

#	ARTICLE	IF	CITATIONS
1	<i>iso</i> -combretastatins A versus Combretastatins A: The Forgotten <i>iso</i> -CA-4 Isomer as a Highly Promising Cytotoxic and Antitubulin Agent. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 4538-4542.	2.9	231
2	New Novobiocin Analogues as Antiproliferative Agents in Breast Cancer Cells and Potential Inhibitors of Heat Shock Protein 90. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 6189-6200.	2.9	136
3	2-Aminobiphenyl Palladacycles: The "Most Powerful" Precatalysts in C-C and C-Heteroatom Cross-Couplings. <i>ACS Catalysis</i> , 2015, 5, 1386-1396.	5.5	136
4	Discovery of Isoerianin Analogues as Promising Anticancer Agents. <i>ChemMedChem</i> , 2011, 6, 488-497.	1.6	128
5	Pd-Catalyzed Reaction of Sterically Hindered Hydrazones with Aryl Halides: Synthesis of Tetra-Substituted Olefins Related to <i>iso</i> -Combretastatin A4. <i>Organic Letters</i> , 2010, 12, 4042-4045.	2.4	111
6	Synthesis and antitumor activity of benzils related to combretastatin A-4. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 3266-3271.	1.0	96
7	DMSO-PdI ₂ as a powerful oxidizing couple of alkynes into benzils: one-pot synthesis of nitrogen-containing five- or six-membered heterocycles. <i>Tetrahedron</i> , 2008, 64, 4287-4294.	1.0	92
8	Synthesis, Biological Evaluation of 1,1-Diarylethylenes as a Novel Class of Antimitotic Agents. <i>ChemMedChem</i> , 2009, 4, 1912-1924.	1.6	82
9	Copper-catalyzed reductive coupling of tosylhydrazones with amines: A convenient route to β -branched amines. <i>Organic and Biomolecular Chemistry</i> , 2011, 9, 6200.	1.5	82
10	Synthesis and biological activity of simplified denoviose-coumarins related to novobiocin as potent inhibitors of heat-shock protein 90 (hsp90). <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 2495-2498.	1.0	80
11	<i>p</i> -Toluenesulfonic acid-promoted selective functionalization of unsymmetrical arylalkynes: a regioselective access to various arylketones and heterocycles. <i>Tetrahedron</i> , 2010, 66, 3775-3787.	1.0	76
12	Platinum Oxide Catalyzed Hydrosilylation of Unsymmetrical Internal Aryl Alkynes under Ortho-Substituent Regiocontrol. <i>Organic Letters</i> , 2005, 7, 5625-5628.	2.4	73
13	One-pot hydrosilylation-protodesilylation of functionalized diarylalkynes: a highly selective access to <i>Z</i> -stilbenes. Application to the synthesis of combretastatin A-4. <i>Tetrahedron Letters</i> , 2008, 49, 1107-1110.	0.7	67
14	Therapeutic Modalities of Squalenoyl Nanocomposites in Colon Cancer: An Ongoing Search for Improved Efficacy. <i>ACS Nano</i> , 2014, 8, 2018-2032.	7.3	67
15	Stereoretentive Palladium-Catalyzed Arylation, Alkenylation, and Alkynylation of α -Thiosugars and Thiols Using Aminobiphenyl Palladacycle Precatalyst at Room Temperature. <i>Chemistry - A European Journal</i> , 2015, 21, 8375-8379.	1.7	66
16	Microwave-assisted efficient synthesis of 1,2-diaryldiketones: a novel oxidation reaction of diarylalkynes with DMSO promoted by FeBr ₃ . <i>Tetrahedron</i> , 2006, 62, 7667-7673.	1.0	65
17	Design, synthesis and anticancer properties of IsoCombretaQuinolines as potent tubulin assembly inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2017, 127, 1025-1034.	2.6	65
18	Conformationally restricted naphthalene derivatives type isocombretastatin A-4 and isoerianin analogues: Synthesis, cytotoxicity and antitubulin activity. <i>European Journal of Medicinal Chemistry</i> , 2012, 52, 22-32.	2.6	64

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19	A convenient and expeditious synthesis of 3-(N-substituted) aminocoumarins via palladium-catalyzed Buchwald-Hartwig coupling reaction. <i>Tetrahedron Letters</i> , 2007, 48, 6928-6932.	0.7	63
20	Nickel-Catalyzed Arylation, Alkenylation, and Alkynylation of Unprotected Thioglycosides at Room Temperature. <i>Chemistry - A European Journal</i> , 2013, 19, 15276-15280.	1.7	60
21	Stereoselective Palladium-Catalyzed Alkenylation and Alkynylation of Thioglycosides. <i>Advanced Synthesis and Catalysis</i> , 2013, 355, 2627-2636.	2.1	59
22	Antiproliferative and apoptotic activities of tosylcyclonovobiocic acids as potent heat shock protein 90 inhibitors in human cancer cells. <i>Cancer Letters</i> , 2009, 274, 88-94.	3.2	57
23	<i>ortho</i> -Substituents Direct Regioselective Addition of Tributyltin Hydride to Unsymmetrical Diaryl (or Heteroaryl) Alkynes: An Efficient Route to Stannylated Stilbene Derivatives. <i>Angewandte Chemie - International Edition</i> , 2002, 41, 1578-1580.	7.2	56
24	Rapid access to 3-(N-substituted)-aminoquinolin-2(1H)-ones using palladium-catalyzed C-N bond coupling reaction. <i>Tetrahedron</i> , 2007, 63, 10202-10210.	1.0	55
25	Synthesis, biological evaluation, and structure-activity relationships of tri- and tetrasubstituted olefins related to isocombretastatin A-4 as new tubulin inhibitors. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 430-442.	1.5	55
26	Design and Synthesis of Tubulin and Histone Deacetylase Inhibitor Based on <i>iso</i> -Combretastatin A-4. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 6574-6591.	2.9	55
27	Palladium-Catalyzed Markovnikov Terminal Arylalkynes Hydrostannation: Application to the Synthesis of 1,1-Diarylethylenes. <i>Journal of Organic Chemistry</i> , 2009, 74, 1337-1340.	1.7	54
28	<i>iso</i> -Combretastatin Quinazolines: Potent Cytotoxic Agents with Antitubulin Activity. <i>ChemMedChem</i> , 2015, 10, 1392-1402.	1.6	52
29	Palladium-Catalyzed Chemoselective and Biocompatible Functionalization of Cysteine-Containing Molecules at Room Temperature. <i>Chemistry - A European Journal</i> , 2016, 22, 11365-11370.	1.7	51
30	Synthesis of substituted quinolines by iron-catalyzed coupling reactions between chloroenynes and Grignard reagents. <i>Tetrahedron Letters</i> , 2004, 45, 1881-1884.	0.7	47
31	Synthetic approach to enyne and enediyne analogues of anticancer agents. <i>Tetrahedron Letters</i> , 2005, 46, 8547-8550.	0.7	47
32	Rapid microwave assisted hydration of internal arylalkynes in the presence of PTSA: an efficient regioselective access to carbonyl compounds. <i>Tetrahedron Letters</i> , 2006, 47, 5497-5501.	0.7	46
33	Copper Acetoacetate [Cu(acac) ₂]/BINAP-Promoted C(sp ³) ₂ -N Bond Formation via Reductive Coupling of N-Tosylhydrazones with Anilines. <i>Advanced Synthesis and Catalysis</i> , 2013, 355, 2417-2429.	2.1	45
34	Tandem One-Pot Palladium-Catalyzed Coupling of Hydrazones, Haloindoles, and Amines: Synthesis of Amino-N-vinylindoles and Their Effect on Human Colon Carcinoma Cells. <i>Journal of Organic Chemistry</i> , 2014, 79, 7583-7592.	1.7	45
35	Stereoretentive Copper-Catalyzed Directed Thioglycosylation of C(sp ²)-H Bonds of Benzamides. <i>Chemistry - A European Journal</i> , 2016, 22, 15006-15010.	1.7	45
36	Efficient Buchwald-Hartwig-Migita Cross-Coupling for DNA Thioglycoconjugation. <i>Chemistry - A European Journal</i> , 2018, 24, 1795-1800.	1.7	45

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37	Bâ€Ringâ€Modified <i>iso</i> Combretastatin Aâ€4 Analogues Endowed with Interesting Anticancer Activities. <i>ChemMedChem</i> , 2011, 6, 2179-2191.	1.6	44
38	Catalytic Three-Component One-Pot Reaction of Hydrazones, Dihaloarenes, and Amines. <i>Organic Letters</i> , 2013, 15, 148-151.	2.4	44
39	Synthesis of (1-â€2)-S-Linked Saccharides and S-Linked Glycoconjugates via a Palladium-G3-XantPhos Precatalyst Catalysis. <i>Journal of Organic Chemistry</i> , 2017, 82, 6720-6728.	1.7	43
40	Palladium(II)-Catalyzed Diastereoselective 2,3-Trans C(sp ³)â€H Arylation of Glycosides. <i>ACS Catalysis</i> , 2018, 8, 7781-7786.	5.5	43
41	1,1-Diheterocyclic Ethylenes Derived from Quinaldine and Carbazole as New Tubulin-Polymerization Inhibitors: Synthesis, Metabolism, and Biological Evaluation. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 1902-1916.	2.9	43
42	Diastereoselective Pd-Catalyzed Anomeric C(sp ³)â€H Activation: Synthesis of Î±-(Hetero)aryl C-Glycosides. <i>ACS Catalysis</i> , 2021, 11, 1818-1826.	5.5	43
43	Stereoselective copper-catalyzed Chanâ€Lamâ€Evans N-arylation of glucosamines with arylboronic acids at room temperature. <i>Chemical Communications</i> , 2013, 49, 8359.	2.2	42
44	Discovery and Biological Activity of 6BrCaQ as an Inhibitor of the Hsp90 Protein Folding Machinery. <i>ChemMedChem</i> , 2011, 6, 804-815.	1.6	40
45	Synthesis of 1,1-Diarylethylenes via Efficient Iron/Copper Co-Catalyzed Coupling of 1-Arylvinyl Halides with Grignard Reagents. <i>Organic Letters</i> , 2012, 14, 2782-2785.	2.4	39
46	Design, synthesis and anticancer properties of 5-arylbenzoxepins as conformationally restricted iso combretastatin A-4 analogs. <i>European Journal of Medicinal Chemistry</i> , 2013, 62, 28-39.	2.6	39
47	Discovery of azaisoerianin derivatives as potential antitumor agents. <i>European Journal of Medicinal Chemistry</i> , 2014, 78, 178-189.	2.6	38
48	Copper-Catalyzed Coupling of <i>N</i> -Tosylhydrazones with Amines: Synthesis of Fluorene Derivatives. <i>ACS Catalysis</i> , 2014, 4, 4498-4503.	5.5	37
49	Xphos ligand and platinum catalysts: A versatile catalyst for the synthesis of functionalized Î²-(E)-vinylsilanes from terminal alkynes. <i>Journal of Organometallic Chemistry</i> , 2008, 693, 2789-2797.	0.8	36
50	Palladium-catalyzed coupling of <i>N</i> -tosylhydrazones with ortho substituted aryl halides: synthesis of 4-arylchromenes and related heterocycles. <i>Tetrahedron Letters</i> , 2011, 52, 1036-1040.	0.7	36
51	Disproportionation reaction of diarylmethylisopropyl ethers: a versatile access to diarylmethanes from diarylcarbinols speeded up by the use of microwave irradiation. <i>Tetrahedron</i> , 2006, 62, 11994-12002.	1.0	34
52	Heat-shock protein 90 inhibitors as antitumor agents: a survey of the literature from 2005 to 2010. <i>Expert Opinion on Therapeutic Patents</i> , 2011, 21, 1501-1542.	2.4	34
53	Synthesis of aryl-thioglycopeptides through chemoselective Pd-mediated conjugation. <i>Chemical Science</i> , 2018, 9, 8753-8759.	3.7	34
54	Palladiumâ€Catalyzed Coupling of 3â€Haloâ€Substituted Coumarins, Chromenes, and Quinolones with Various Nitrogenâ€Containing Nucleophiles. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 5077-5088.	1.2	33

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55	Developments of isoCombretastatin A-4 derivatives as highly cytotoxic agents. <i>European Journal of Medicinal Chemistry</i> , 2020, 190, 112110.	2.6	33
56	Tosylcyclonovobiocic acids promote cleavage of the hsp90-associated cochaperone p23. <i>Biochemical and Biophysical Research Communications</i> , 2009, 379, 514-518.	1.0	32
57	Ni/Photoredox-Dual-Catalyzed Functionalization of 1-Thiosugars. <i>Organic Letters</i> , 2019, 21, 5132-5137.	2.4	32
58	Rapid synthesis of 4-arylchromenes from ortho-substituted alkynols: A versatile access to restricted isocombretastatin A-4 analogues as antitumor agents. <i>European Journal of Medicinal Chemistry</i> , 2015, 90, 834-844.	2.6	31
59	Recent Advances in Transition-Metal-Catalyzed Functionalization of 1-Thiosugars. <i>Asian Journal of Organic Chemistry</i> , 2018, 7, 2026-2038.	1.3	31
60	An efficient coupling of N-tosylhydrazones with 2-halopyridines: synthesis of 2- <i>alk-1-en-1-yl</i> pyridines endowed with antitumor activity. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 3664.	1.5	30
61	Palladium-Catalyzed One-Pot Reaction of Hydrazones, Dihaloarenes, and Organoboron Reagents: Synthesis and Cytotoxic Activity of 1,1-Diarylethylene Derivatives. <i>Journal of Organic Chemistry</i> , 2015, 80, 6715-6727.	1.7	28
62	Recent advances in the synthesis of pyrido[1,2- <i>cd</i>]indoles. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 3509-3526.	1.5	27
63	Synthesis and antiproliferative activity of novobiocin analogues as potential hsp90 inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2014, 83, 498-507.	2.6	26
64	Electrochemical nickel-catalyzed Migita cross-coupling of 1-thiosugars with aryl, alkenyl and alkynyl bromides. <i>Chemical Communications</i> , 2020, 56, 4464-4467.	2.2	25
65	Regioselective hydrostannation of diarylalkynes directed by a labile ortho bromine atom: An easy access to stereodefined triarylolefins, hybrids of combretastatin A-4 and isocombretastatin A-4. <i>European Journal of Medicinal Chemistry</i> , 2010, 45, 3617-3626.	2.6	24
66	One-Pot Assembly of Unsymmetrical Biaryl Thioglycosides through Chemoselective Palladium-Catalyzed Three-Component Tandem Reaction. <i>Organic Letters</i> , 2018, 20, 4067-4071.	2.4	23
67	N,N-bis-heteroaryl methylamines: Potent anti-mitotic and highly cytotoxic agents. <i>European Journal of Medicinal Chemistry</i> , 2019, 168, 176-188.	2.6	23
68	Formulation and in vitro efficacy of liposomes containing the Hsp90 inhibitor 6BrCaQ in prostate cancer cells. <i>International Journal of Pharmaceutics</i> , 2016, 499, 101-109.	2.6	20
69	Palladium-Catalyzed One-Pot Synthesis of 5-(1-Arylvinyl)-1H-benzimidazoles: Overcoming the Limitation of Acetamide Partners. <i>Advanced Synthesis and Catalysis</i> , 2016, 358, 1833-1847.	2.1	19
70	Intramolecular Pd-Catalyzed Arylation of 1-Amidosugars: A New Route to N-Glycosyl Quinolin-2-ones. <i>Organic Letters</i> , 2016, 18, 2126-2129.	2.4	19
71	Suzuki Coupling Reactions of <i>E</i> - and <i>Z</i> -Chloroenynes with Boronic Acids: Versatile Access to Functionalized 1,3-Enynes. <i>European Journal of Organic Chemistry</i> , 2010, 2010, 725-731.	1.2	18
72	A palladium-catalyzed coupling of 3-chloroquinoxalinones with various nitrogen-containing nucleophiles. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 3808.	1.5	18

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73	Synthesis of 2-substituted indoles through cyclization and demethylation of 2-alkynyl dimethylanilines by ethanol. <i>Green Chemistry</i> , 2019, 21, 4204-4210.	4.6	18
74	Anticancer properties of indole derivatives as IsoCombretastatin A-4 analogues. <i>European Journal of Medicinal Chemistry</i> , 2021, 223, 113656.	2.6	18
75	Identification of a new series of flavopiridol-like structures as kinase inhibitors with high cytotoxic potency. <i>European Journal of Medicinal Chemistry</i> , 2020, 199, 112355.	2.6	17
76	Discovery and Hit to Lead Optimization of Novel Combretastatin A-4 Analogues: Dependence of C-Linker Length and Hybridization. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2013, 13, 1614-1635.	0.9	17
77	Cyclic bridged analogs of isoCA-4: Design, synthesis and biological evaluation. <i>European Journal of Medicinal Chemistry</i> , 2021, 209, 112873.	2.6	16
78	The Metabolic Fate of <i>iso</i> Combretastatin A-4 in Human Liver Microsomes: Identification, Synthesis and Biological Evaluation of Metabolites. <i>ChemMedChem</i> , 2011, 6, 1781-1788.	1.6	15
79	PtO ₂ /PTSA system catalyzed regioselective hydration of internal arylalkynes bearing electron withdrawing groups. <i>RSC Advances</i> , 2018, 8, 11536-11542.	1.7	15
80	Copper-Catalyzed Anomeric O-Arylation of Carbohydrate Derivatives at Room Temperature. <i>Journal of Organic Chemistry</i> , 2019, 84, 9226-9238.	1.7	15
81	Room-Temperature Pd-Catalyzed Synthesis of 1-(Hetero)aryl Selenoglycosides. <i>Organic Letters</i> , 2020, 22, 6584-6589.	2.4	14
82	Synthesis and Biological Activities of Pyrazino[1,2-a]indole and Pyrazino[1,2-a]indol-1-one Derivatives. <i>Pharmaceuticals</i> , 2021, 14, 779.	1.7	14
83	Antitumor activity of nanoliposomes encapsulating the novobiocin analog 6BrCaQ in a triple-negative breast cancer model in mice. <i>Cancer Letters</i> , 2018, 432, 103-111.	3.2	13
84	Diversity-oriented synthesis of fused thioglycosyl benzo[e][1,4]oxathiepin-5-ones and benzo[f][1,4]thiazepin-5(2H)-ones by a sequence of palladium-catalyzed glycosyl thiol arylation and deprotection-lactonization reactions. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 10904-10916.	1.5	12
85	Synthesis of 2,3-Substituted ² N-Glycosyl Indoles through C-H Activation/Annulation Process under Rh(III)-Catalysis. <i>Organic Letters</i> , 2020, 22, 57-61.	2.4	12
86	Design, synthesis and biological evaluation of quinoline-2-carbonitrile-based hydroxamic acids as dual tubulin polymerization and histone deacetylases inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2022, 240, 114573.	2.6	12
87	Regiochemical Aspects of the Platinum Oxide Catalyzed Hydrosilylation of Alkynes. <i>Synthesis</i> , 2007, 2007, 2025-2036.	1.2	11
88	Selective Palladium-Catalyzed Domino Heck/Buchwald-Hartwig Arylations of <i>N</i> -Glycosylcinnamamides: An Efficient Route to 4-Aryl- <i>N</i> -Glycosylquinolin-2-ones. <i>Advanced Synthesis and Catalysis</i> , 2017, 359, 1320-1330.	2.1	11
89	Toward a Greener Barluenga-Cross-Coupling: Microwave-Promoted C-C Bond Formation with a Pd/PEG/H ₂ O Recyclable Catalytic System. <i>Organic Letters</i> , 2019, 21, 8708-8712.	2.4	11
90	Ortho-Substituents Direct Regioselective Addition of Tributyltin Hydride to Unsymmetrical Diaryl (or) Triarylmethyl Ketones. <i>Journal of Organic Chemistry</i> , 2011, 76, 1648-1650.	1.6	10

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91	Regio- and diastereoselective Pd-catalyzed synthesis of C2-aryl glycosides. <i>Chemical Communications</i> , 2020, 56, 7175-7178.	2.2	10
92	A general Pd/Cu-catalyzed C-H heteroarylation of 3-bromoquinolin-2(1H)-ones. <i>Organic and Biomolecular Chemistry</i> , 2014, 12, 8533-8541.	1.5	9
93	One-Pot Synthesis of 2-Styrylindoles from <i>Ortho</i> -Substituted Chloroenynes. <i>Journal of Organic Chemistry</i> , 2018, 83, 15323-15332.	1.7	8
94	Convergent Synthesis of N,S-bis Glycosylquinolin-2-ones via a Pd-G3-XantPhos Precatalyst Catalysis. <i>Molecules</i> , 2018, 23, 519.	1.7	8
95	Reversing Reactivity: Stereoselective Desulfurative 1,2- <i>trans</i> -O-Glycosylation of Anomeric Thiosugars with Carboxylic Acids under Copper or Cobalt Catalysis. <i>Journal of Organic Chemistry</i> , 2020, 85, 8893-8909.	1.7	7
96	A Convenient Metal-Free Synthesis of (E)-3-Styrylisocoumarins through Annulation of (E)-1,4-Diarylenynes. <i>Synthesis</i> , 2016, 48, 3382-3392.	1.2	6
97	Synthesis and functionalization of 3-bromo-2-(2-chlorovinyl)benzothiophenes as molecular tools. <i>RSC Advances</i> , 2017, 7, 46007-46013.	1.7	6
98	Mild Deprotection of Dithioacetals by TMSCl/Nal Association in CH ₃ CN. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 5775-5779.	1.2	6
99	Synthesis of N-Glycosyl-2-oxindoles by Pd-Catalyzed N-Arylation of 1-Amidosugars. <i>Organic Letters</i> , 2020, 22, 4201-4206.	2.4	6
100	Synthesis of <i>Trifluoromethyl Arylsulfoximine Thioglycosides</i> through Pd-Catalyzed Migita Cross-Coupling. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 4972-4981.	1.2	6
101	Synthesis and Biological Activity of 3-(Heteroaryl)quinolin-2(1H)-ones Bis-Heterocycles as Potential Inhibitors of the Protein Folding Machinery Hsp90. <i>Molecules</i> , 2022, 27, 412.	1.7	6
102	Controllable Activation of Pd-G3 Palladacycle Precatalyst in the Presence of Thiosugars: Rapid Access to <i>Aminobiphenyl Thioglycoside Atropoisomers</i> at Room Temperature. <i>Chemistry - an Asian Journal</i> , 2017, 12, 3114-3118.	1.7	4
103	Unexpected Oxidative Ring Opening of Electron-Rich 3-Aminobenzofurans into \pm -Ketoimines Derivatives. <i>Journal of Organic Chemistry</i> , 2019, 84, 1725-1733.	1.7	4
104	Synthesis and antiproliferative activity of 6BrCaQ-TPP conjugates for targeting the mitochondrial heat shock protein TRAP1. <i>European Journal of Medicinal Chemistry</i> , 2022, 229, 114052.	2.6	4
105	Azoliums and Ag(I)-Heterocyclic Carbene Thioglycosides: Synthesis, Reactivity and Bioactivity. <i>European Journal of Organic Chemistry</i> , 2022, 2022, .	1.2	4
106	Copper-catalyzed sulfonylation of <i>N</i> -tosylhydrazones followed by a one-pot C-N bond formation. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 5358-5367.	1.5	3
107	Synthesis and Anticancer Properties of Oxazepines Related to Azaisoerianin and IsoCoQuines. <i>ChemMedChem</i> , 2020, 15, 1571-1578.	1.6	2
108	Synthesis of axially chiral biaryl thioglycosides through thiosugar-directed Pd-catalyzed asymmetric C-H activation. <i>Chemical Communications</i> , 2021, 57, 10355-10358.	2.2	2