

Sylvie Radix

List of Publications by Year in descending order

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#	ARTICLE	IF	CITATIONS
1	A Journey through <i>Hemetsberger</i> – <i>Knittel</i> , <i>Leimgruber</i> – <i>Batcho</i> and <i>Bartoli</i> Reactions: Access to Several Hydroxy 5- and 6-Azaindoles. <i>Helvetica Chimica Acta</i> , 2022, 105, .	1.6	1
2	Solvent- and metal-free hydroboration of alkynes under microwave irradiation. <i>Tetrahedron Letters</i> , 2020, 61, 151596.	1.4	9
3	Spectral data for the synthesis of (E)-alkenylboronic acid pinacol esters via hydroboration of alkynes. <i>Data in Brief</i> , 2020, 30, 105354.	1.0	0
4	Synthesis and biological evaluation of zinc chelating compounds as metallo- β -lactamase inhibitors. <i>MedChemComm</i> , 2019, 10, 528-537.	3.4	13
5	Synthesis and biological evaluation of new dipicolylamine zinc chelators as metallo- β -lactamase inhibitors. <i>Tetrahedron</i> , 2019, 75, 1525-1540.	1.9	10
6	N,N ϵ -2-disubstituted cinnamamide derivatives potentiate ciprofloxacin activity against overexpressing NorA efflux pump <i>Staphylococcus aureus</i> 1199B strains. <i>European Journal of Medicinal Chemistry</i> , 2018, 150, 900-907.	5.5	12
7	Enhancement of iodinin solubility by encapsulation into cyclodextrin nanoparticles. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018, 33, 370-375.	5.2	7
8	Synthesis and Preclinical Evaluation of TPA-Based Zinc Chelators as Metallo- β -lactamase Inhibitors. <i>ACS Infectious Diseases</i> , 2018, 4, 1407-1422.	3.8	35
9	Deoxycholic acid derivatives as inhibitors of P-glycoprotein-mediated multidrug efflux. <i>Steroids</i> , 2016, 116, 5-12.	1.8	9
10	Unsymmetrical β -end-functionalized oligo(cyclohexylidenes): efficient synthesis and conformational analysis. <i>Tetrahedron</i> , 2016, 72, 4032-4038.	1.9	2
11	Progesterone–adenine hybrids as bivalent inhibitors of P-glycoprotein-mediated multidrug efflux: Design, synthesis, characterization and biological evaluation. <i>Steroids</i> , 2012, 77, 1177-1191.	1.8	8
12	Design, synthesis and evaluation of progesterone–adenine hybrids as bivalent inhibitors of P-glycoprotein-mediated multidrug efflux. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 3165-3168.	2.2	6
13	Synthesis and modulation properties of imidazo[4,5-b]pyridin-7-one and indazole-4,7-dione derivatives towards the <i>Cryptosporidium parvum</i> CpABC3 transporter. <i>European Journal of Medicinal Chemistry</i> , 2010, 45, 2480-2488.	5.5	6
14	Regioselective N-alkylation of imidazo[4,5-b]pyridine-4-oxide derivatives: an experimental and DFT study. <i>Tetrahedron Letters</i> , 2009, 50, 1828-1833.	1.4	12
15	A Simple and High-Yield Route to N-9-purinyl-acetic Acid Derivatives Coupled with Amino Acids. <i>Letters in Organic Chemistry</i> , 2008, 5, 484-489.	0.5	2
16	Total synthesis of two natural phenanthrenes: confusarin and a regioisomer. <i>Tetrahedron</i> , 2007, 63, 12379-12387.	1.9	20
17	Synthesis of [6,n] cis-fused ring compounds via Cr-mediated dearomatisation–ring-closing metathesis. <i>Organic and Biomolecular Chemistry</i> , 2006, 4, 342-351.	2.8	14
18	A Regioselective Route to 5- and 6-Azaindoles. <i>Synlett</i> , 2005, 2005, 2080-2082.	1.8	0

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19	Efficient Access to Fused Ring Compounds via Dearomatization/Ring-Closing Metathesis.. ChemInform, 2004, 35, no.	0.0	0
20	Efficient Access to Fused Ring Compounds via Dearomatization/Ring-Closing Metathesis. Synlett, 2003, 2003, 2407-2409.	1.8	12